

TITLE OF THE INVENTION

**ELECTROPHOTOGRAPHIC PHOTORECEPTOR, AND IMAGE FORMING METHOD,
IMAGE FORMING APPARATUS AND PROCESS CARTRIDGE FOR IMAGE FORMING
APPARATUS USING THE ELECTROPHOTOGRAPHIC PHOTORECEPTOR**

5

BACKGROUND OF THE INVENTION

Field of the Invention

The present invention relates to an electrophotographic
photoreceptor, and an image forming method, an image forming
10 apparatus and process cartridge for image forming apparatus using
the electrophotographic photoreceptor.

Discussion of the Background

Recently, information-processing systems using an
15 electrophotographic method are making a remarkable progress.
In particular, laser printers and digital copiers which record
data with light by changing the data into digital signals make
remarkable improvements in their printing qualities and
reliabilities. Further, technologies used in these printers
20 and copiers are applied to laser printers and digital copiers
capable of printing full-color images with high-speed printing
technologies. Because of these reasons, photoreceptors are
required both to produce high-quality images and to have high
durability.

25 Photoreceptors using organic photosensitive materials
are widely used for these laser printers and digital copiers
due to their cost, productivity and non-polluting properties.

The organic photoreceptors are generally classified to a single-layered type and a functionally-separated type. The first practical organic photoreceptor, i.e., PVK-TNF charge transfer complex photoreceptor was the former single-layered type. In 1968, Mr. Hayashi and Mr. Regensburger independently invented PVK/a-Se multi-layered photoreceptor. In 1977, Mr. Melz, and in 1978, Mr. Schlosser disclosed a multi-layered photoreceptor whose photosensitive layers are all formed from organic materials, i.e., an organic-pigment dispersed layer and an organic low-molecular-weight material dispersed polymer layer. These are called as functionally-separated photoreceptors because of having a charge generation layer (CGL) generating a charge by absorbing light and a charge transport layer (CTL) transporting the charge and neutralizing the charge on a surface of the photoreceptor. The multi-layered photoreceptor has much more improved sensitivity and durability than the single-layered photoreceptor. In addition, since materials can be separately selected for a charge generation material (CGM) and a charge transport material (CTM), a choice range of the materials is largely expanded. Because of these reasons, the multi-layered photoreceptor is now prevailing in the market.

A mechanism to form an electrostatic latent image in the multi-layered photoreceptor is as follows:

the photoreceptor is charged and irradiated with light; the light passes through the CTL and is absorbed by the CGM in the CGL to generate a charge; the charge is injected into the

CTL at an interface of the CGL and the CTL; and the charge moves in the CTL by an electric field and neutralizes the charge on the surface of the photoreceptor to form an electrostatic latent image.

5 However, the photosensitive layers of the organic photoreceptor are easily abraded due to repeated use, and therefore potential and photosensitivity of the photoreceptor tend to deteriorate, resulting in background fouling due to a scratch on the surface thereof and deterioration of density and
10 quality of the resultant images. Therefore, abrasion resistance of the organic photoreceptor has been an important subject. Further, recently, in accordance with speeding up of the printing speed and downsizing of an image forming apparatus, the photoreceptor has to have a smaller diameter, and durability
15 thereof becomes a more important subject.

 As a method of improving the abrasion resistance of the photoreceptor, methods of imparting lubricity to the photosensitive layer, hardening the photosensitive layer, including a filler therein and using a high-molecular-weight
20 CTM instead of a low-molecular-weight CTM are widely known. However, another problem occurs when these methods are used to prevent the abrasion of the photoreceptor. Namely, an oxidized gas such as ozone and NO_x arising due to use conditions or environment, adheres to the surface of the photosensitive layer
25 and decreases the surface resistance thereof, resulting in a problem such as blurring of the resultant images. So far, such a problem has been avoided to some extent because the material

causing the blurred images are gradually scraped off in accordance with the abrasion of the photosensitive layer. However, in order to comply with the above-mentioned recent demand for higher sensitivity and durability of the photoreceptor, a new technique has to be imparted thereto. In order to decrease an influence of the material causing the blurred images, there is a method of equipping the photoreceptor with a heater, which is a large drawback for downsizing the apparatus and decreasing the electric power consumption. In addition, a method of including an additive such as an antioxidant in the photosensitive layer is effective, but since a simple additive does not have photoconductivity, including much amount thereof in the photosensitive layer causes problems such as deterioration of the sensitivity and increase of residual potential of the resultant photoreceptor.

As mentioned above, the electrophotographic photoreceptor having less abrasion by being imparted with abrasion resistance or a process design around thereof inevitably produces blurred and low-resolution images, and it is difficult to have both of high durability and high quality of the resultant images. This is because high surface resistance of the photosensitive layer is preferable to prevent the blurred images and low surface resistance thereof is preferable to prevent the increase of residual potential.

Japanese Laid-Open Patent Publication No. 2000-231204 or 2002-313111 discloses a method of including at least a compound having a dialkylamino group in a photosensitive layer to solve

the above-mentioned problem such as blurring of the resultant images due to a blur generating material such as an oxidizing gas. The reason why the compound is effective for maintaining quality of the resultant images after repeated use is not clarified at this time. However, it is supposed that the dialkylamino group having a strong basic neutralizes the oxidizing gas which is considered to cause the blurred images. However, the compound has an effect on image quality after the repeated use, but the resultant photoreceptor does not have high sensitivity and cannot comply with high speed printing because of having a low charge transportability. Therefore, an addition amount thereof has a limit, and a method of combining the compound with a CTM to increase sensitivity and repeated use stability of the resultant photoreceptor is disclosed therein.

On the other hand, it is described that a stilbene compound having a dialkylamino group disclosed in Japanese Laid-Open Patent Publication No. 60-196768 and Japanese Patent No. 2884353 has an effect on the blurred images due to the oxidizing gas on page 37 of Konica Technical Report Vol. 13 written by Itami, etc. and published in 2000. However, since the compound has a substituted dialkylamino group having a strong mesomeric effect (+M effect) at a resonance portion in its triarylamine structure, which is a charge transport site, total ionization potential is extremely small. Therefore, the compound has a critical defect of being quite difficult to practically use because charge retainability of a photosensitive layer in which the compound is used alone as a CTM largely deteriorates from the beginning

or after repeated use. In addition, even when the above-mentioned stilbene compound is used together with other CTMs as it is in the present invention, the compound has a considerably smaller ionization potential than the other CTMs and becomes a trap site against a charge transport, and therefore, the resultant photoreceptor has quite a low sensitivity and a large residual potential.

Because of these reasons, a need exists for an electrophotographic photoreceptor having high durability against repeated use for a long time, preventing deterioration of image density and blurred images and stably producing quality images.

SUMMARY OF THE INVENTION

Accordingly, an object of the present invention is to provide an electrophotographic photoreceptor having high durability against repeated use for a long time, preventing deterioration of image density and blurred images and stably producing high-quality images.

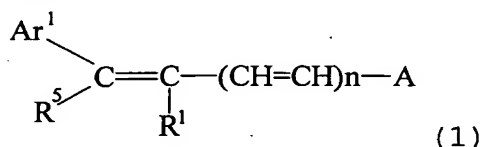
Another object of the present invention is to provide an image forming method, an image forming apparatus and a process cartridge using the photoreceptor, in which the photoreceptor need not be exchanged, which enables downsizing the apparatus in accordance with the high-speed printing or smaller diameter of the photoreceptor, and which stably produce high-quality images even after repeated use for a long time.

Briefly these objects and other objects of the present

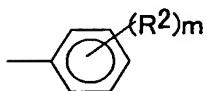
invention as hereinafter will become more readily apparent can be attained by an electrophotographic photoreceptor including an electroconductive substrate and a photosensitive layer on the electroconductive substrate, wherein the photosensitive layer includes at least a compound having a substituted or unsubstituted alkylamino group and a charge transport material, and wherein an oxidation potential (Eox1) of the substituted or unsubstituted alkylamino group and an oxidation potential (Eox2) of the charge transport material satisfy the following relationship (I):

$$E_{ox1} - E_{ox2} \geq -0.2 \quad (I)$$

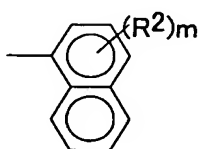
The charge transport material is preferably a stilbene compound having the following formula (1):



wherein n is 0 or 1; R¹ represents a hydrogen atom, an alkyl group or a substituted or unsubstituted phenyl group; Ar¹ represents a substituted or unsubstituted aryl group; R⁵ represents an alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aryl group; and A represents a 9-anthryl group, a substituted or unsubstituted carbazoly group or a group having the following formula (4) or (5):

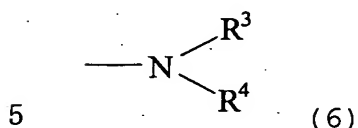


(4),



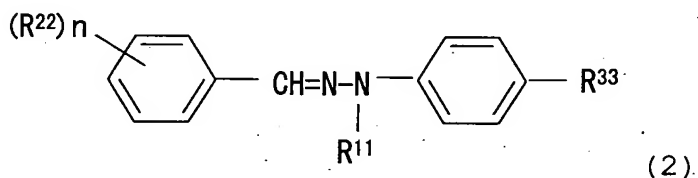
(5)

wherein R^2 represents a hydrogen atom, an alkyl group, an alkoxy group, a halogen atom or a group having the following formula (6); and m is an integer of from 1 to 3;



wherein R^3 and R^4 independently represent a substituted or unsubstituted aromatic ring group, and may form a ring, and wherein R^2 may be the same or different from each other when m is not less than 2, and A and R^1 may form a ring together when n is 0.

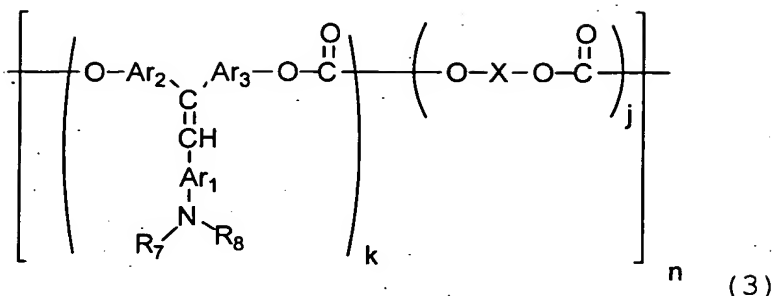
Further, the charge transport material is preferably a hydrazone compound having the following formula (2):



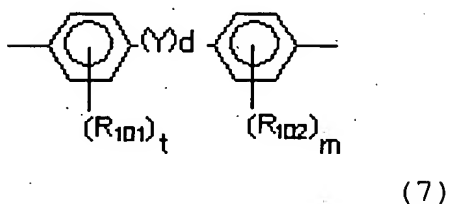
15 wherein the R^{11} represents an alkyl group, a benzyl group, a phenyl group or a naphthyl group; R^{22} represents a hydrogen atom, an alkyl group having 1 to 3 carbon atoms, an alkoxy group having 1 to 3 carbon atoms, a dialkylamino group, a diaralkylamino group or a substituted or unsubstituted diarylamino group; n represents
20 integers of from 1 to 4 and R^{22} is optionally the same or different from each other when n is not less than 2; and R^{33} represents a hydrogen atom or a methoxy group.

Furthermore, the charge transport polymer material is preferably a charge transport polymer material having the

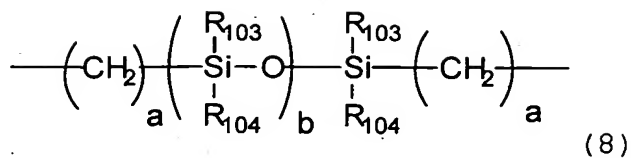
following formula (3):



wherein R^7 and R^8 independently represent a substituted or
 5 unsubstituted aromatic ring group; Ar^1 , Ar^2 and Ar^3 independently
 represent an aromatic ring group; k is a number of from 0.1 to
 1.0 and j is a number of from 0 to 0.9; n represents a repeating
 number and is an integer of from 5 to 5,000; and X represents
 10 a divalent group having the following formula (7):



wherein, R^{101} and R^{102} independently represent a substituted or
 unsubstituted alkyl group, a substituted or unsubstituted aryl
 15 group, or a halogen atom; t and m independently represent 0 or
 an integer of from 1 to 4; d is 0 or 1; and Y represents a linear
 alkylene group, a branched alkylene group, a cyclic alkylene
 group, $-\text{O}-$, $-\text{S}-$, $-\text{SO}-$, $-\text{SO}_2-$, $-\text{CO}-$, $-\text{CO}-\text{O}-\text{Z}-\text{O}-\text{CO}-$ (Z represents
 a divalent aliphatic group), or a group having the following
 20 formula (8):



wherein, a is an integer of from 1 to 20; b is an integer of from 1 to 2,000; and R^{103} and R^{104} independently represent a substituted or unsubstituted alkyl group, or a substituted or unsubstituted aryl group, and wherein R^{101} , R^{102} , R^{103} and R^{104} may be the same or different from the others.

These and other objects, features and advantages of the present invention will become apparent upon consideration of the following description of the preferred embodiments of the present invention taken in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

Various other objects, features and attendant advantages of the present invention will be more fully appreciated as the same becomes better understood from the detailed description when considered in connection with the accompanying drawings in which like reference characters designate like corresponding parts throughout and wherein:

Fig. 1 is a cross-sectional view of an embodiment of the photosensitive layer of the electrophotographic photoreceptor of the present invention;

Fig. 2 is a cross-sectional view of another embodiment

of the photosensitive layer of the electrophotographic photoreceptor of the present invention;

Fig. 3 is a cross-sectional view of a third embodiment of the photosensitive layer of the electrophotographic photoreceptor of the present invention;

Fig. 4 is a cross-sectional view of a fourth embodiment of the photosensitive layer of the electrophotographic photoreceptor of the present invention;

Fig. 5 is a cross-sectional view of a fifth embodiment of the photosensitive layer of the electrophotographic photoreceptor of the present invention;

Fig. 6 is a schematic view illustrating a partial cross-section of an embodiment of the electrophotographic image forming apparatus of the present invention;

Fig. 7 is a schematic view for explaining an embodiment of the electrophotographic image forming process of the present invention;

Fig. 8 is a schematic view illustrating a cross-section of an embodiment of the process cartridge of the present invention;

Fig. 9 is a chart showing a relationship between a difference (ΔE) between the oxidation potential of the compound having an alkylamino group and that of a CTM, and a bright section potential (VL) in the electrophotographic photoreceptor of the present invention;

Fig. 10 is a chart showing a relationship between a difference (ΔE) between the oxidation potential of the compound

having an alkylamino group and that of another CTM, and a bright section potential (VL) in the electrophotographic photoreceptor of the present invention;

Fig. 11 is a chart showing a XD spectrum of an
5 oxotitaniumphthalocyanine powder of the present invention;

Fig. 12 is a chart showing a relationship between a difference (ΔE) between the oxidation potential of the compound having an alkylamino group and that of a third CTM, and a bright section potential (VL) in the electrophotographic photoreceptor
10 of the present invention;

Fig. 13 is a chart showing a XD spectrum of another oxotitaniumphthalocyanine powder of the present invention; and

Fig. 14 is a chart showing a relationship between a difference (ΔE) between the oxidation potential of the compound
15 having an alkylamino group and that of a fourth CTM, and a bright section potential (VL) in the electrophotographic photoreceptor of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

20 Generally, the present invention provides an electrophotographic photoreceptor having high durability against repeated use for a long time, preventing deterioration of image density and blurred images and stably producing quality images.

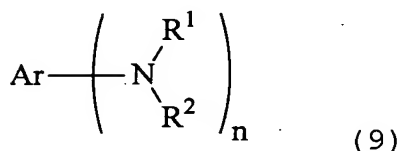
25 The electrophotographic photoreceptor of the present invention includes an electroconductive substrate and a photosensitive layer on the electroconductive substrate,

wherein the photosensitive layer includes at least a compound having a substituted or unsubstituted alkylamino group and a charge transport material, and wherein an oxidation potential (Eox1) of the substituted or unsubstituted alkylamino group and
5 an oxidation potential (Eox2) of the charge transport material satisfy the following relationship (I):

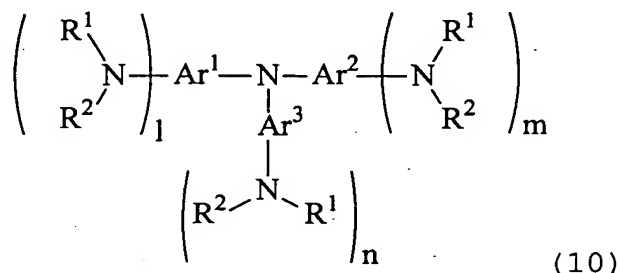
$$E_{ox1} - E_{ox2} \geq -0.2 \quad (I)$$

The electrophotographic photoreceptor including a compound having an alkylamino group which is mixed with a CTM
10 of the present invention has a high sensitivity and a high durability, and stably produces high-quality images even after repeated use when the compound having an alkylamino group has an ionization potential not less than that of CTM by a certain level. Namely, as mentioned above, as the alkylamino group is
15 a substituent having a strong mesomeric effect (+M effect), a total ionization potential of the compound becomes extremely small when the substituent is substituted at a resonance portion thereof. When the compound having an alkylamino group has considerably a smaller ionization potential than that of the
20 CTM, it becomes a hole trap site against the charge transport, and therefore the resultant electrophotographic photoreceptor has quite a low sensitivity and a large residual potential. When the compound having an alkylamino group has an ionization potential not less than that of CTM by a certain level, the
25 resultant electrophotographic photoreceptor has a high sensitivity and a high durability, and stably produces high-quality images even after repeated use.

Specific examples of the compound having an alkylamino group include compounds having the following formulae (9) to (35):

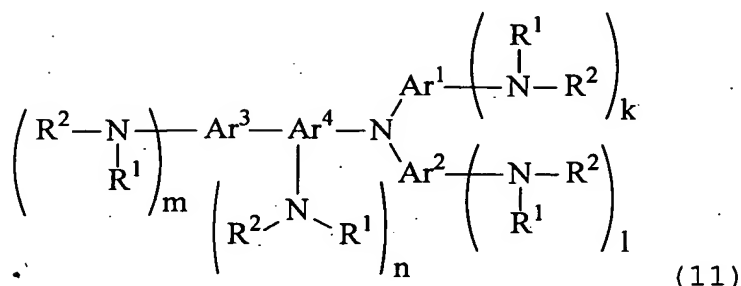


wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; n represents an integer of from 1 to 4; and Ar represents a substituted or unsubstituted aromatic ring group;

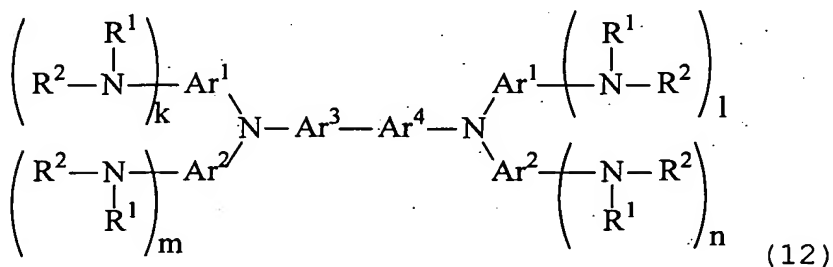


wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; 1 , m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; Ar^1 , Ar^2 and Ar^3 independently represent a substituted or unsubstituted aromatic ring group; and Ar^1 and Ar^2 , Ar^2 and Ar^3

or Ar³ and Ar¹ may independently form a heterocyclic group including a nitrogen atom together;

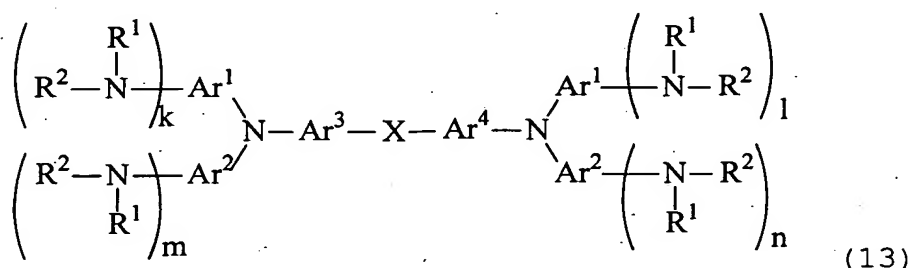


wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; k, l, m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; Ar¹, Ar², Ar³ and Ar⁴ independently represent a substituted or unsubstituted aromatic ring group; and Ar¹ and Ar², Ar¹ and Ar⁴ or Ar³ and Ar⁴ may independently form a ring together;

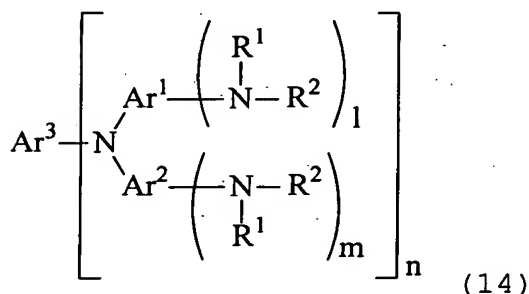


wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group

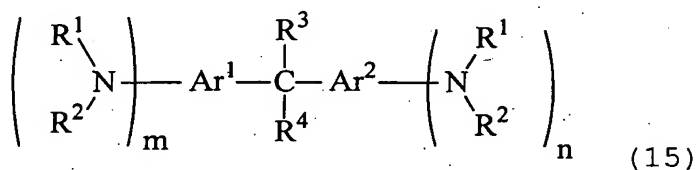
including a nitrogen atom; k, l, m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; Ar¹, Ar², Ar³ and Ar⁴ independently represent a substituted or unsubstituted aromatic ring group; and Ar¹ and Ar², Ar¹ and Ar³ or Ar³ and Ar⁴ may independently form a ring together;



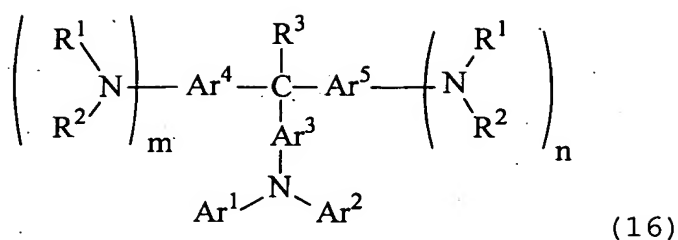
wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; k, l, m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; Ar¹, Ar², Ar³ and Ar⁴ independently represent a substituted or unsubstituted aromatic ring group; Ar¹ and Ar², Ar¹ and Ar³ or Ar¹ and Ar⁴ may independently form a ring together; and X represents a methylene group, a cyclohexylidene group, an oxy atom or a sulfur atom;



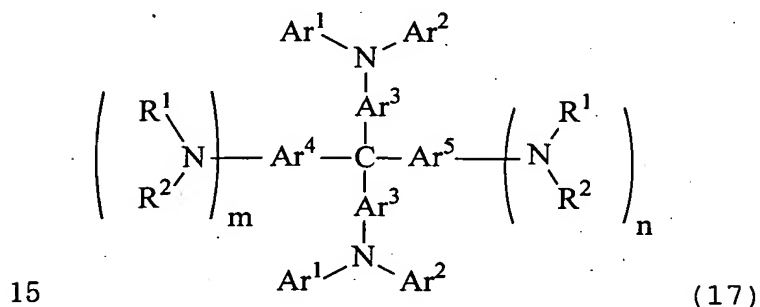
wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; 1 and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; Ar¹, Ar² and Ar³ independently represent a substituted or unsubstituted aromatic ring group; Ar¹ and Ar² or Ar¹ and Ar³ may independently form a ring together; and n represents an integer of from 1 to 4;



wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; R³ and R⁴ independently represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 11 carbon atoms and a substituted or unsubstituted aromatic ring group; and Ar¹ and Ar² independently represent a substituted or unsubstituted aromatic ring group, and one of Ar¹, Ar², R³ and R⁴ is an aromatic heterocyclic group;

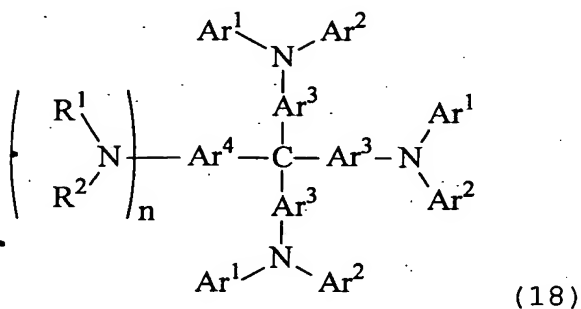


wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; R^3 represents a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 11 carbon atoms and a substituted or unsubstituted aromatic ring group; Ar^1 , Ar^2 , Ar^3 , Ar^4 and Ar^5 independently represent a substituted or unsubstituted aromatic ring group; and Ar^1 and Ar^2 or Ar^1 and Ar^3 may form a heterocyclic group including a nitrogen atom together;

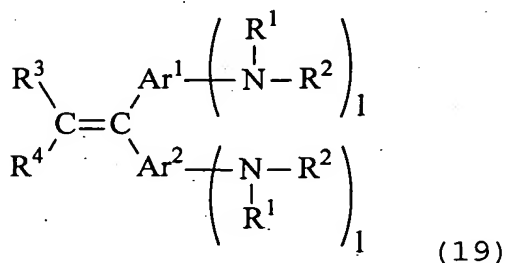


wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; m and n independently represent 0

or an integer of from 1 to 3, and are not 0 at the same time;
 Ar^1 , Ar^2 , Ar^3 , Ar^4 and Ar^5 independently represent a substituted
 or unsubstituted aromatic ring group; and Ar^1 and Ar^2 or Ar^1 and
 Ar^3 may form a heterocyclic group including a nitrogen atom
 5 together;

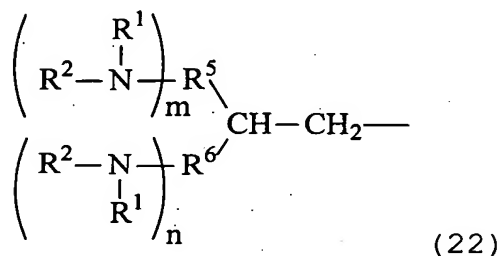


wherein R^1 and R^2 independently represent an alkyl group having
 1 to 4 carbon atoms, which is substituted with an aromatic ring
 10 group or an unsubstituted alkyl group having 1 to 4 carbon atoms
 and may be combined with each other to form a heterocyclic group
 including a nitrogen atom; n represents an integer of from 1
 to 3; Ar^1 , Ar^2 , Ar^3 and Ar^4 independently represent a substituted
 or unsubstituted aromatic ring group; and Ar^1 and Ar^2 or Ar^1 and
 15 Ar^3 may form a heterocyclic group including a nitrogen atom
 together;

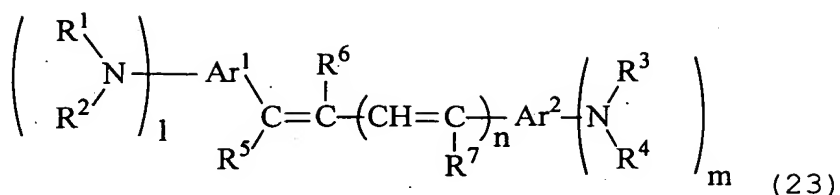


wherein R^1 and R^2 independently represent an alkyl group having
 20 1 to 4 carbon atoms, which is substituted with an aromatic ring

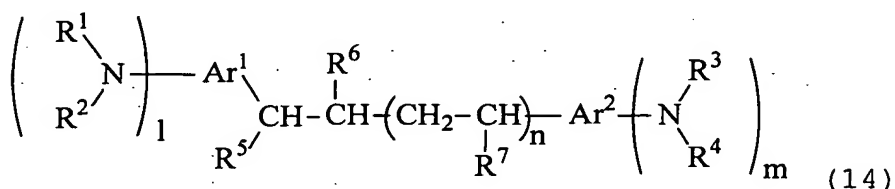
wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; n represents an integer of from 1 to 3; Ar¹ and Ar² independently represent a substituted or unsubstituted aromatic ring group; R³ and R⁴ independently represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms, a substituted or unsubstituted aromatic ring group or a group having the following formula (22), and R³ and R⁴ are not hydrogen atoms at the same time:



wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; m and n independently represent 0 or an integer of from 1 to 3; and R⁵ and R⁶ independently represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group, and wherein R³ and R⁴, R⁵ and R⁶ or Ar¹ and Ar² may independently form a ring together;

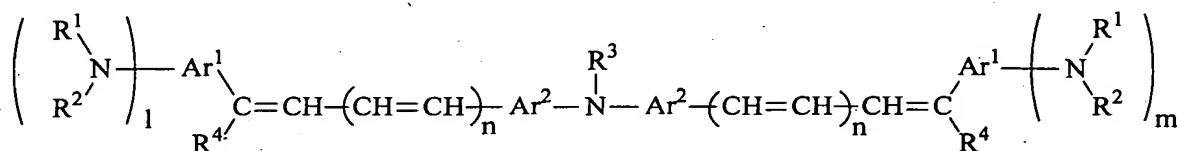


wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; R^3 and R^4 independently represent a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; R^5 , R^6 and R^7 independently represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; Ar^1 and Ar^2 independently represent a substituted or unsubstituted aromatic ring group; R^3 and R^4 or Ar^2 and R^4 may form a heterocyclic group including a nitrogen atom together; Ar^1 and R^5 may form a ring together; 1 represents an integer of from 1 to 3; m represents 0 or an integer of from 1 to 3; and n represents 0 or 1;



wherein R^1 and R^2 independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group

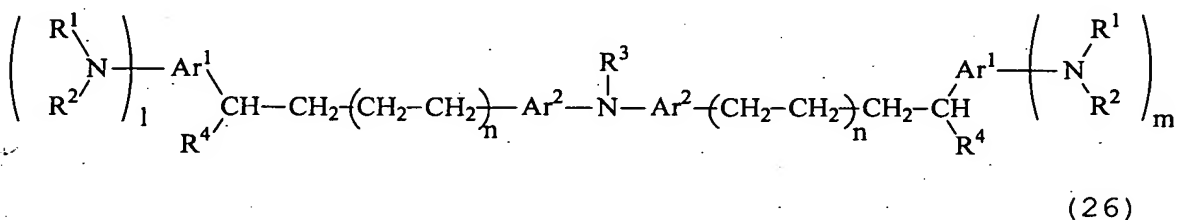
including a nitrogen atom; R³ and R⁴ independently represent a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; R⁵, R⁶ and R⁷ independently represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; Ar¹ and Ar² independently represent a substituted or unsubstituted aromatic ring group; R³ and R⁴ or Ar² and R⁴ may form a heterocyclic group including a nitrogen atom together; Ar¹ and R⁵ may form a ring together; l represents an integer of from 1 to 3; m represents 0 or an integer of from 1 to 3; and n represents 0 or 1;



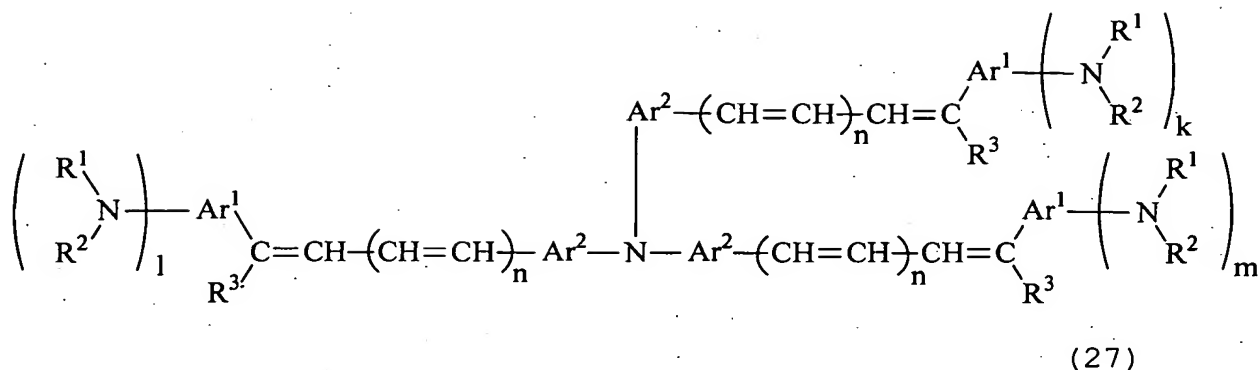
(25)

wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; l and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; R³ represents a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; R⁴ represents a hydrogen atom, a substituted or

unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; Ar¹ and Ar² represent a substituted or unsubstituted aromatic ring group; Ar¹ and R⁴, Ar² and R³ or Ar² and another Ar² may form a ring together; and n represents 0 or 1;

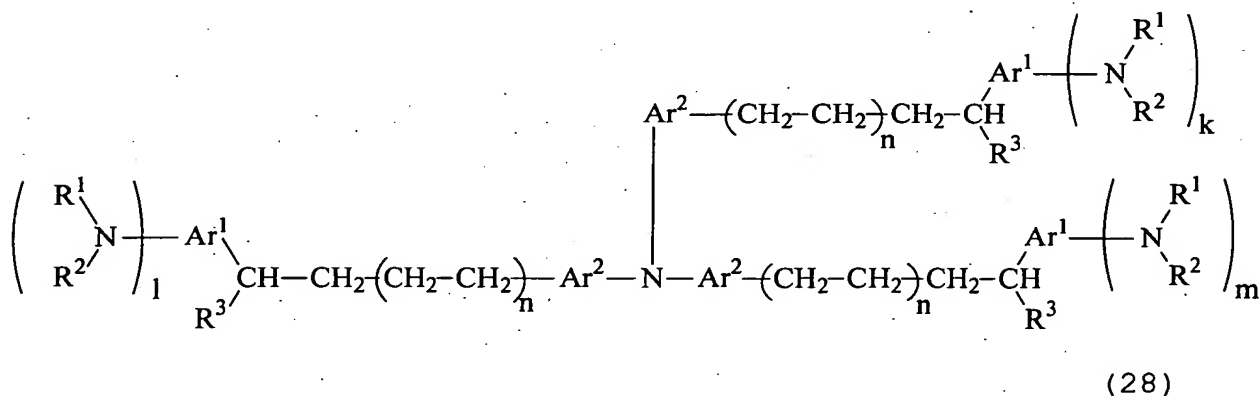


wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; l and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; R³ represents a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; R⁴ represents a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; Ar¹ and Ar² represent a substituted or unsubstituted aromatic ring group; Ar¹ and R⁴, Ar² and R³ or Ar² and another Ar² may form a ring together; and n represents 0 or 1;



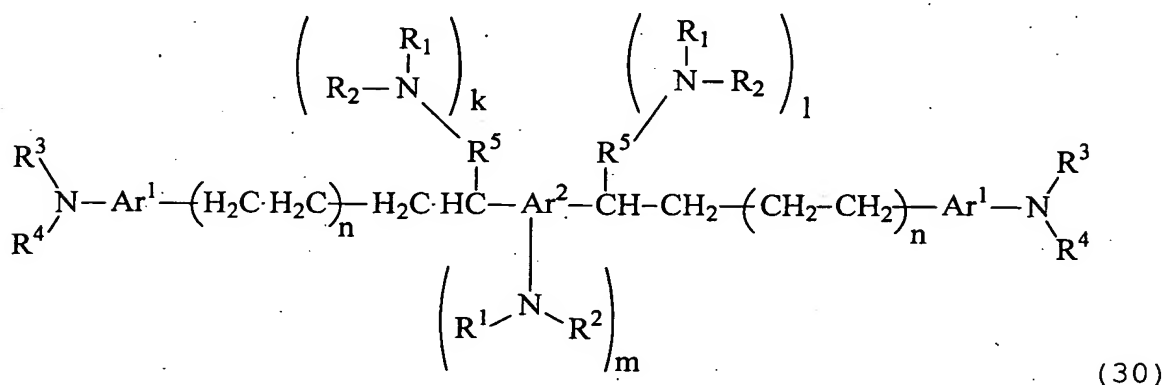
wherein R^1 and R^2 independently represent an alkyl group having
 1 to 4 carbon atoms, which is substituted with an aromatic ring
 5 group or an unsubstituted alkyl group having 1 to 4 carbon atoms
 and may be combined with each other to form a heterocyclic group
 including a nitrogen atom; k , l and m independently represent
 0 or an integer of from 1 to 3, and are not 0 at the same time;
 R^3 represents a hydrogen atom, a substituted or unsubstituted
 10 alkyl group having 1 to 4 carbon atoms or a substituted or
 unsubstituted aromatic ring group; Ar^1 and Ar^2 represent a
 substituted or unsubstituted aromatic ring group; Ar^1 and R^4 ,
 Ar^2 and R^3 or Ar^2 and another Ar^2 may form a ring together; and
 n represents 0 or 1;

15



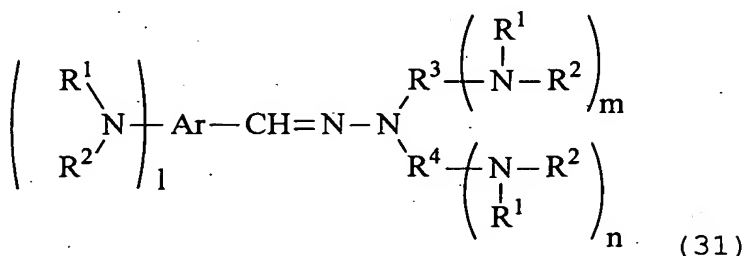
wherein R^1 and R^2 independently represent an alkyl group having

unsubstituted aromatic ring group; Ar¹ and Ar² represent a
 substituted or unsubstituted aromatic ring group; R³ and R⁴ or
 Ar¹ and R⁴ may form a heterocyclic group including a nitrogen
 atom together; k, l and m independently represent 0 or an integer
 5 of from 1 to 3; n represents 1 or 2; and R³ and R⁴ independently
 represent an alkyl group having 1 to 4 carbon atoms and may be
 combined with each other to form a heterocyclic group including
 a nitrogen atom when k, l and m are 0 at the same time;

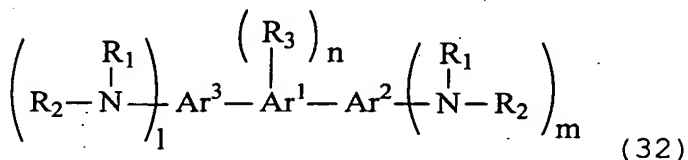


wherein R¹ and R² independently represent an alkyl group having
 1 to 4 carbon atoms, which is substituted with an aromatic ring
 group or an unsubstituted alkyl group having 1 to 4 carbon atoms
 and may be combined with each other to form a heterocyclic group
 including a nitrogen atom; R³ and R⁴ independently represent
 15 a substituted or unsubstituted alkyl group having 1 to 4 carbon
 atoms or a substituted or unsubstituted aromatic ring group;
 R⁵ represents a hydrogen atom, a substituted or unsubstituted
 alkyl group having 1 to 4 carbon atoms or a substituted or
 20 unsubstituted aromatic ring group; Ar¹ and Ar² represent a
 substituted or unsubstituted aromatic ring group; R³ and R⁴ or
 Ar¹ and R⁴ may form a heterocyclic group including a nitrogen

atom together; m represents 0 or an integer of from 1 to 4; n represents 1 or 2; and R³ and R⁴ independently represent an alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom when m is 0;

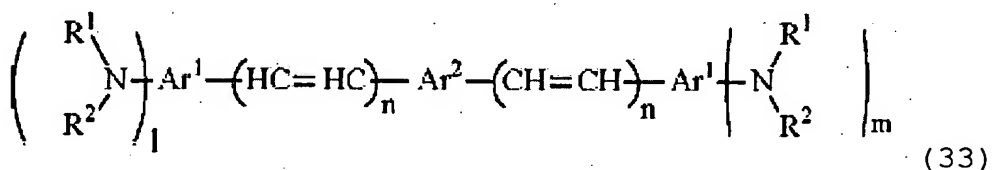


wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms and may be combined with each other to form a heterocyclic group including a nitrogen atom; Ar represents a substituted or unsubstituted aromatic ring group; R³ and R⁴ represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; and 1, m and n independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time;

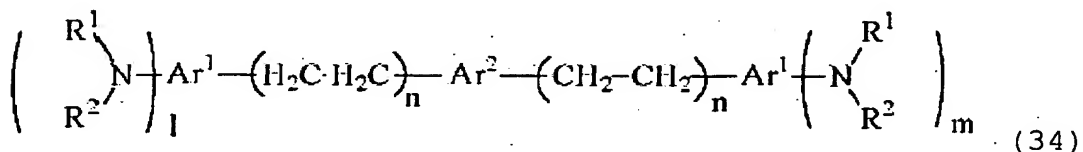


wherein R¹ and R² independently represent an alkyl group having 1 to 4 carbon atoms, which is substituted with an aromatic ring group or an unsubstituted alkyl group having 1 to 4 carbon atoms

and may be combined with each other to form a heterocyclic group including a nitrogen atom; Ar¹, Ar² and Ar³ represent a substituted or unsubstituted aromatic ring group; R³ represents a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aromatic ring group; l and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; and n represents an integer of from 1 to 3;

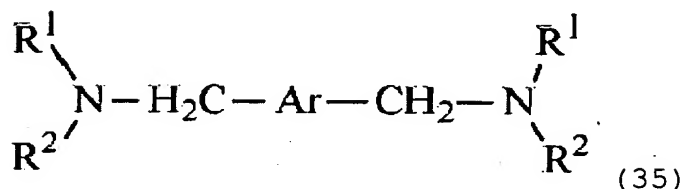


wherein R¹ and R² independently represent an alkyl group substituted with an aromatic hydrocarbon group or an unsubstituted alkyl group and may be combined with each other to form a heterocyclic group including a nitrogen atom; Ar¹ and Ar² represent a substituted or unsubstituted aromatic ring group; l and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; and n represents 1 or 2;



wherein R¹ and R² independently represent an alkyl group substituted with an aromatic hydrocarbon group or an unsubstituted alkyl group and may be combined with each other to form a heterocyclic group including a nitrogen atom; Ar¹ and Ar² represent a substituted or unsubstituted aromatic ring group;

1 and m independently represent 0 or an integer of from 1 to 3, and are not 0 at the same time; and n represents 1 or 2; and



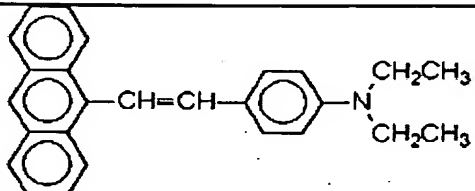
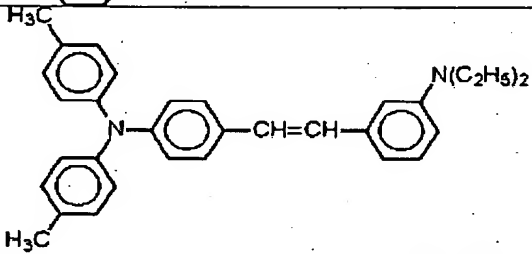
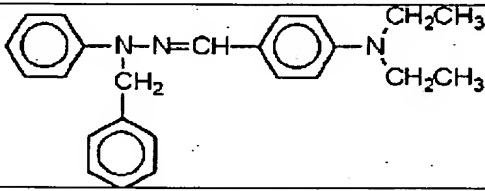
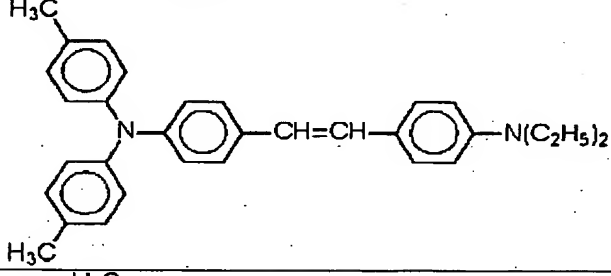
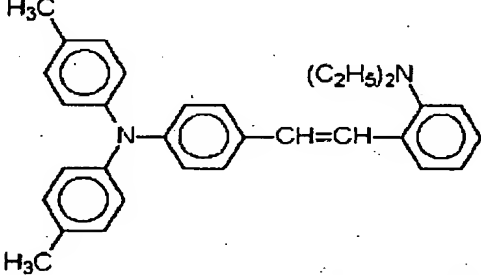
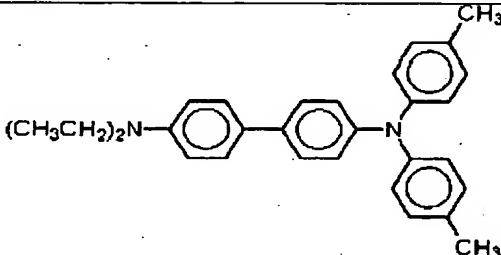
5 wherein R¹ and R² independently represent substituted or unsubstituted alkyl group and a substituted or unsubstituted aromatic hydrocarbon group, and one of R¹ and R² is a substituted or unsubstituted aromatic hydrocarbon group, and may be combined with each other to form a heterocyclic group including a nitrogen
10 atom; and Ar represents a substituted or unsubstituted aromatic hydrocarbon group.

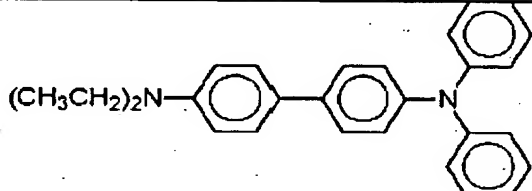
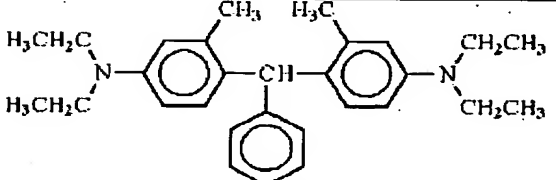
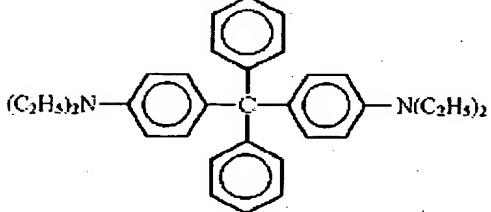
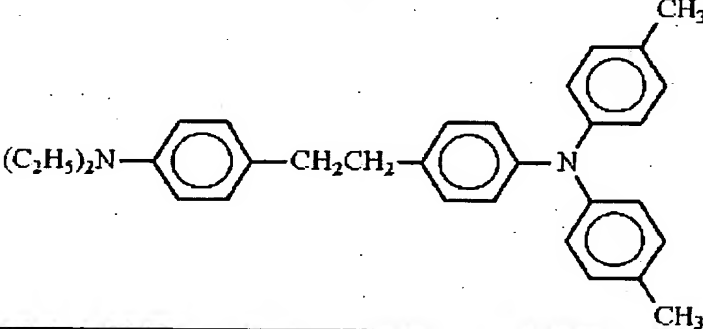
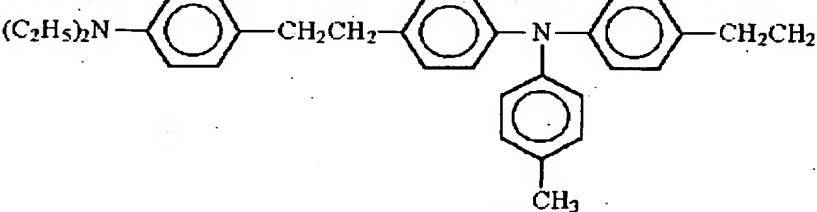
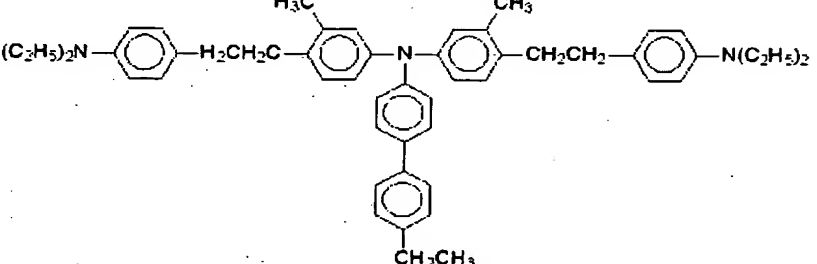
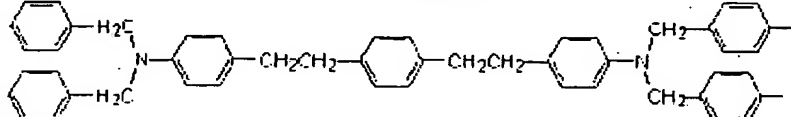
Specific examples of the alkyl group mentioned in the explanations of these formulae (9) to (35) include a methyl group, an ethyl group, a propyl group, a butyl group, a hexyl group,
15 an undecanyl group, etc. Specific examples of the aromatic hydrocarbon group include aromatic ring groups such as benzene, biphenyl, naphthalene, anthracene, fluorene and pyrene; and aromatic heterocyclic groups such as pyridine, quinoline, thiophene, furan, oxazole, oxadiazole and carbazole. Specific
20 examples of their substituents include the above-mentioned specific examples of the alkyl group; an alkoxy group such as a methoxy group, an ethoxy group, a propoxy group and a butoxy group; a halogen atoms such as a fluorine atom, a chlorine atom, a bromine atom and an iodine atom; the above-mentioned aromatic
25 hydrocarbon groups; and heterocyclic ring groups such as

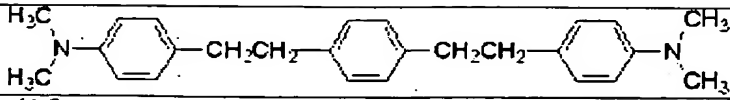
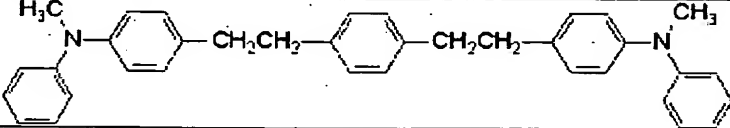
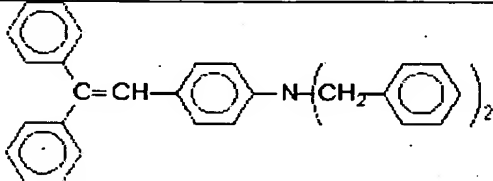
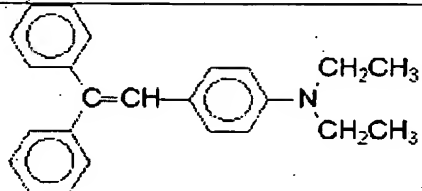
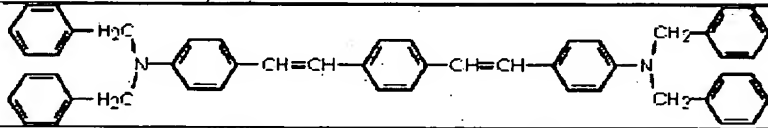
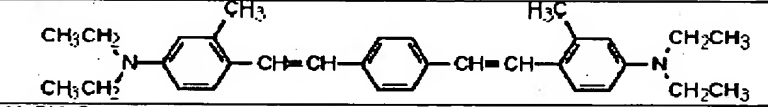
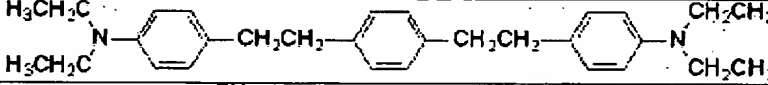
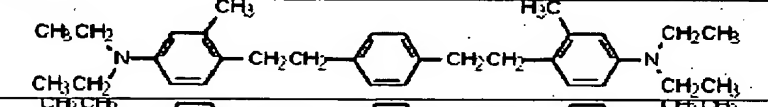

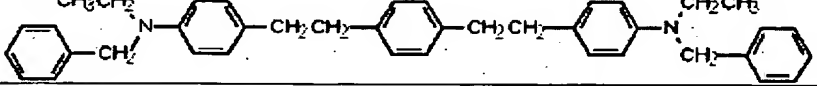

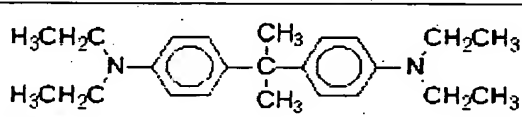
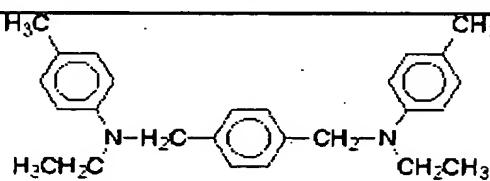
pyrrolidine, piperidine and piperazine. When R^1 and R^2 are combined with each other to form a heterocyclic group including a nitrogen atom, specific examples thereof include a condensed heterocyclic group such as pyrrolidino groups, piperidino groups and piperazino groups condensed with aromatic hydrocarbon groups.

A method of measuring the oxidation potential (E_{ox1}) of the compound having an alkylamino group and that (E_{ox2}) of the CTM, i.e., a primary oxidation half-wave potential. A predetermined amount of acetonitrile and that of an unrelated salt (a supporting electrolyte) such as tetrabutylammonium perchlorate and tetraethylammonium perchlorate are added to a material to be measured to prepare a test liquid. The oxidation potential of the material can be measured by analyzing the test liquid with an electrochemical analysis methods such as polarographic methods and cyclic voltammetric methods. The electrochemical analysis methods are disclosed in "Electrochemical Methods" written by A. J. Bard and L. R. Faulkner and published by Wiley in 1980 in detail, wherein a potential scanning method using a potentiostat is used, and wherein a dropping mercury electrode is used as a working electrode, a noble metal such as platinum or gold (platinum in "Electrochemical Methods") as a counter electrode and a saturated calomel electrode (SCE) as a reference electrode.

Specific examples of the compound having an alkylamino group and oxidation potential (E_{ox1}) thereof are shown as follows:

Compound No.	Formula	Eox (V vs. SCE)
1		0.520
2		0.605
3		0.500
4		0.440
5		0.655
6		0.520

7		0.550
8		0.650
9		0.640
10		0.660
11		0.660
12		0.625
13		0.875

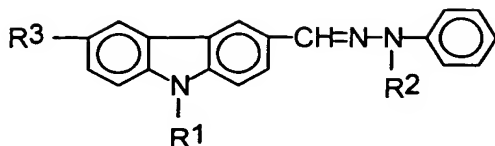
14		0.660
15		0.780
16		0.750
17		0.600
18		0.625
19		0.395
20		0.660
21		0.620
22		0.545
23		0.760
24		0.460
25		0.785
26		0.740

27		0.750
28		0.600
29		0.855
30		0.680
31		0.620
32		0.750
33		0.780
34		0.770

Hereinafter, the CTM will be explained. The CTM is classified to a low-molecular-weight CTM and a charge transport polymer material.

Specific examples of the low-molecular-weight CTM include compounds having the following formulae (1), (2), (4) to (6),

and (36) to (54).

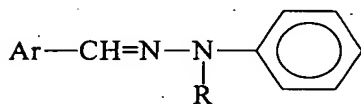


(36)

wherein R¹ represents a methyl group, an ethyl group, a
2-hydroxyethyl group or a 2-chlorethyl group; and R² represents
a methyl group, an ethyl group, a benzyl group or a phenyl group;
and R³ represents a hydrogen atom, a chlorine atom, a bromine
atom, an alkyl group having 1 to 4 carbon atoms, an alkoxy group
having 1 to 4 carbon atoms, a dialkylamino group or a nitro group.

Specific examples of the compound having formula (36)
include

9-ethylcalbazole-3-aldehyde-1-methyl-1-phenylhydrazone,
9-ethylcalbazole-3-aldehyde-1-benzyl-1-phenylhydrazone,
9-ethylcalbazole-3-aldehyde-1,1-diphenylhydrazone, etc.

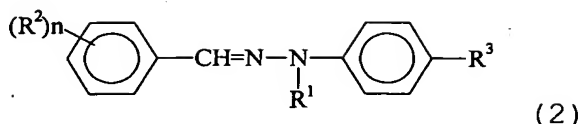


(37)

wherein Ar represents a naphthalene ring, an anthracene ring,
a pyrene ring and their substituents, a pyridine ring, a furan
ring or thiophene ring; and R represents an alkyl group, a phenyl
group or a benzyl group.

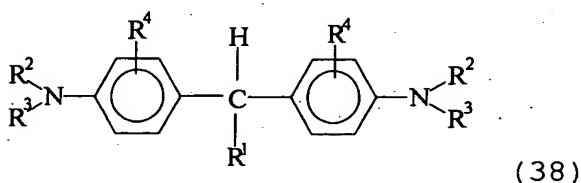
Specific examples of the compound having formula (37)
include 4-diethylaminostyryl- β -
-aldehyde-1-methyl-1-phenylhydrazone,
4-methoxynaphthalene-1-aldehyde-1-benzyl-1-phenylhydrazone,

etc.



wherein R^1 represents an alkyl group, a benzyl group, a phenyl group or a naphthyl group; R^2 represents a hydrogen atom, an alkyl group having 1 to 3 carbon atoms, an alkoxy group having 1 to 3 carbon atoms, a dialkylamino group, diaralkylamino group or a diarylamino group; n represents an integer of from 1 to 4 and R^2 may be the same or different from each other when n is not less than 2; and R^3 represents a hydrogen atom or a methoxy group.

Specific examples of the compound having formula (2) include 4-methoxybenzaldehyde-1-methyl-1-phenylhydrazone, 2,4-dimethoxybenzaldehyde-1-benzyl-1-phenylhydrazone, 4-diethylaminobenzaldehyde-1,1-diphenylhydrazone, 4-methoxybenzaldehyde-1-(4-methoxy)phenylhydrazone, 4-diphenylaminobenzaldehyde-1-benzyl-1-phenylhydrazone, 4-dibenzylaminobenzaldehyde-1,1-diphenylhydrazone, etc.

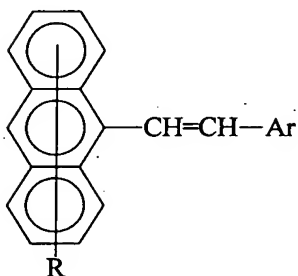


wherein R^1 represents an alkyl group having 1 to 11 carbon atoms, a substituted or unsubstituted phenyl group or a heterocyclic ring group; R^2 and R^3 independently represent a hydrogen atom, an alkyl group having 1 to 4 carbon atoms, a hydroxyalkyl group, a chloralkyl group or a substituted or unsubstituted aralkyl

group, and may be combined each other to form a heterocyclic ring group including a nitrogen atom; and R^4 independently represent a hydrogen atom, an alkyl group having 1 to 4 carbon atoms, an alkoxy group or a halogen atom.

- 5 Specific examples of the compound having the formula (38) include 1,1-bis(4-dibenzylaminophenyl)propane, tris(4-diethylaminophenyl)methane, 1,1-bis(4-dibenzylaminophenyl)propane, 2,2'-dimethyl-4,4'-bis(diethylamino)-triphenylmethane, etc.

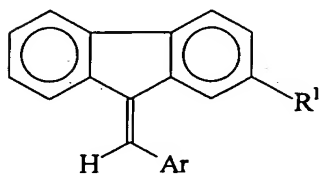
10



(39)

wherein R represents a hydrogen atom or a halogen atom; and Ar represents a substituted or unsubstituted phenyl group, a naphthyl group, an anthryl group or a carbazolyl group.

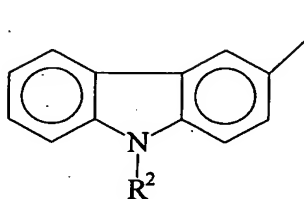
- 15 Specific examples of the compound having the formula (39) include 9-(4-diethylaminostyryl)anthracene, 9-bromo-10-(4-diethylaminostyryl)anthracene, etc.



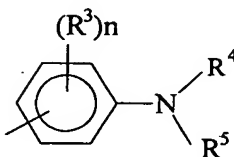
(40)

- 20 wherein R^1 represents a hydrogen atom, a cyano group, an alkoxy group having 1 to 4 carbon atoms or a alkyl group having 1 to

4 carbon atoms; and Ar represents a group having the following formulae (41) and (42):



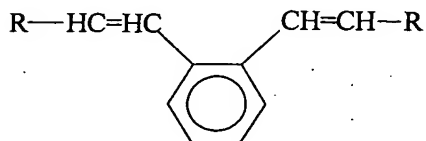
(41)



(42)

wherein R^2 represents an alkyl group having 1 to 4 carbon atoms; R^3 represents a hydrogen atom, a halogen atom, an alkyl group having 1 to 4 carbon atoms, an alkoxy group having 1 to 4 carbon atoms or a dialkylamino group; n is 1 or 2, and R^3 may be the same or different from each other when n is 2; and R^4 and R^5 represent a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted benzyl group.

Specific examples of the compound having the formula (40) include 9-(4-dimethylaminobenzylidene)fluorene, 3-(9-fluorenylidene)-9-ethylcarbazole, etc.

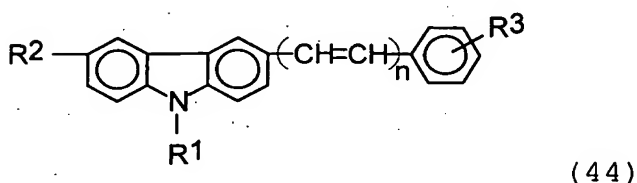


(43)

wherein R represents a carbazolyl group, a pyridyl group, a thienyl group, an indolyl group, a furyl group, a substituted or unsubstituted phenyl, styryl, naphthyl group or an anthryl group, and their substituents are selected from the group consisting of a dialkylamino group, an alkyl group, an alkoxy

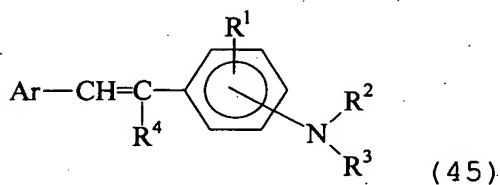
group, a carboxyl group or its ester, a halogen atom, a cyano group, an aralkylamino group, N-alkyl-N-aralkylamino group, an amino group, a nitro group and an acethylamino group.

Specific examples of the compound having the formula (43) include 1,2-bis-(4-diethylaminostyryl)benzene,
1,2-bis(2-,4-dimethoxystyryl)benzene, etc.



wherein R^1 represents a lower alkyl group, a substituted or unsubstituted phenyl group or a benzyl group; R^2 and R^3 represent a hydrogen atom, a lower alkyl group, a lower alkoxy group, a halogen atom, a nitro group, an amino group or an amino group substituted by a lower alkyl group or a benzyl group; and n is 1 or 2.

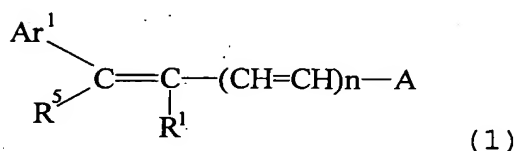
Specific examples of the compound having the formula (44) include 3-styryl-9-ethylcarbazole,
3-(4-methoxystyryl)-9-ethylcarbazole, etc.



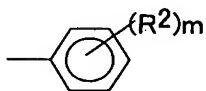
wherein R^1 represents a hydrogen atom, an alkyl group, an alkoxy group or a halogen atom; R^2 and R^3 represent a substituted or unsubstituted aryl group; R^4 represents a hydrogen atom, a lower alkyl group or a substituted or unsubstituted phenyl group; and

Ar represents a substituted or unsubstituted phenyl group or a naphthyl group.

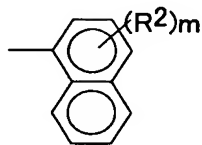
Specific examples of the compound having the formula (31) include 4-diphenylaminostilbene, 4-dibenzylaminostilbene, 4-ditolylaminostilbene, 1-(4-phenylaminostyryl)naphthalene, 1-(4-diethylaminostyryl)naphthalene, etc.



wherein n is 0 or 1; R¹ represents a hydrogen atom, an alkyl group or a substituted or unsubstituted phenyl group; Ar¹ represents a substituted or unsubstituted aryl group; R⁵ represents an alkyl group having 1 to 4 carbon atoms or a substituted or unsubstituted aryl group; and A represents a 9-anthryl group, a substituted or unsubstituted carbazoly group or a group having the following formula (4) or (5):

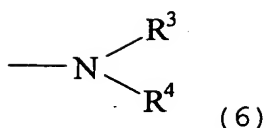


(4)



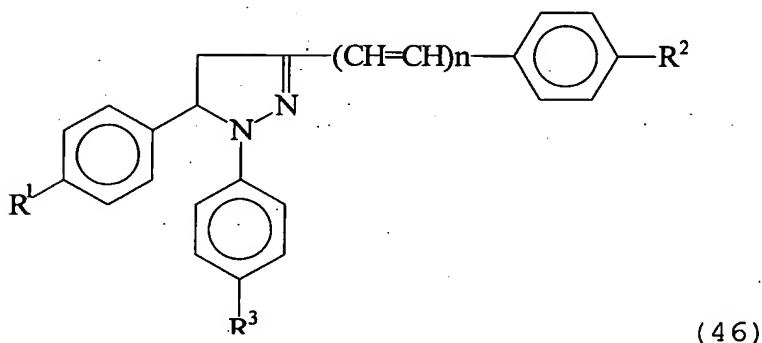
(5)

wherein R² represents a hydrogen atom, an alkyl group, an alkoxy group, a halogen atom or a group having the following formula; and m is an integer of from 1 to 3;



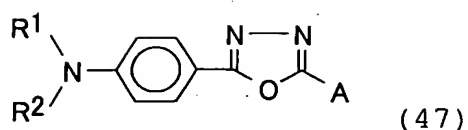
wherein R^3 and R^4 independently represent a substituted or unsubstituted aryl group, and R^4 may form a ring, and wherein R^2 may be the same or different from each other when m is not less than 2, and A and R^1 may form a ring together when n is 0.

Specific examples of the compound having the formula (1) include 4'-diphenylamino- α -phenylstilbene, 4'-bis(4-methylphenyl)amino- α -phenylstilbene, etc.



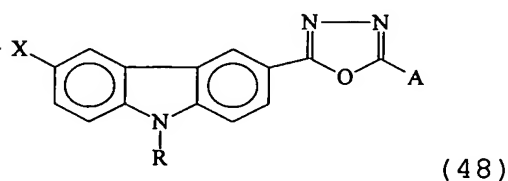
wherein R^1 , R^2 and R^3 represent a hydrogen atom, a lower alkyl group, a lower alkoxy group, a halogen atom or a dialkylamino group; and n is 0 or 1.

Specific examples of the compound having the formula (46) include 1-phenyl-3-(4-diethylaminostyryl)-5-(4-diethylaminophenyl)pyrazoline, etc.



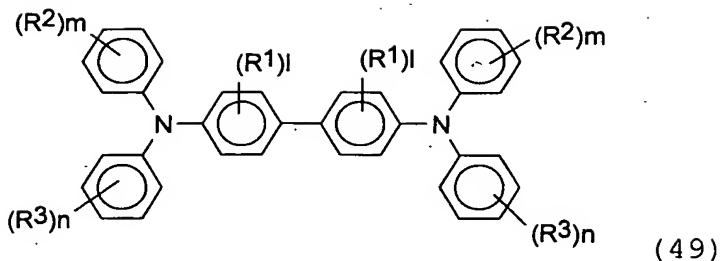
wherein R¹ and R² represent an alkyl group including a substituted alkyl group or a substituted or unsubstituted aryl group; and A represents a substituted amino group, a substituted or
 5 unsubstituted aryl group or an aryl group.

Specific examples of the compound having the formula (47) include 2,5-bis(4-diethylaminophenyl)-1,3,4-oxadiazole, 2-N,N-diphenylamino-5-(4-diethylaminophenyl)-1,3,4-oxadiazole,
 10 2-(4-dimethylaminophenyl)-5-(4-diethylaminophenyl)-1,3,4-oxadiazole, etc.



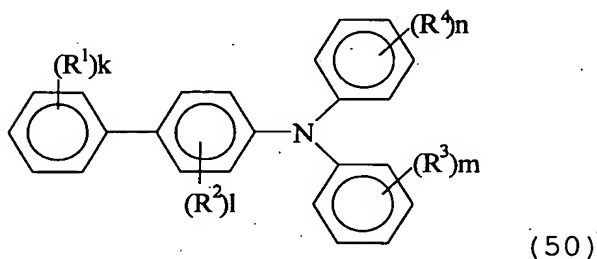
wherein X represents a hydrogen atom, a lower alkyl group or
 15 a halogen atom; R represents an alkyl group including a substituted alkyl group or a substituted or unsubstituted aryl group; and A represents a substituted amino group, a substituted or unsubstituted aryl group or an aryl group.

Specific examples of the compound having the formula (48) include
 20 2-N,N-diphenylamino-5-(N-ethylcarbazole-3-yl)-1,3,4-oxadiazole,
 2-(4-diethylaminophenyl)-5-(N-ethylcarbazole-3-yl)-1,3,4-oxadiazole, etc.



wherein R^1 represents a lower alkyl group, a lower alkoxy group or a halogen atom; R^2 and R^3 independently represent a hydrogen atom, a lower alkyl group, a lower alkoxy group or a halogen atom; and l , m and n independently represent 0 or an integer of from 1 to 4.

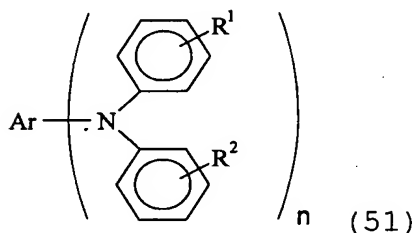
Specific examples of the benzidine compound having the formula (49) include N,N' -diphenyl- N,N' -bis(3-methylphenyl)-[1,1'-biphenyl]-4,4'-diamine, 3,3'-dimethyl- N,N,N',N' -tetrakis(4-methylphenyl)-[1,1'-biphenyl]-4,4'-diamine, etc.



wherein R^1 , R^3 and R^4 represent a hydrogen atom, an amino group, an alkoxy group, a thioalkoxy group, an aryloxy group, a methylenedioxy group, a substituted or unsubstituted alkyl group, a halogen atom or a substituted or unsubstituted aryl group; R^2 represents a hydrogen atom, an alkoxy group, a substituted or unsubstituted alkyl group or a halogen atom, but a case in

which R^1 , R^2 , R^3 and R^4 are all hydrogen atoms is excluded; and k , l , m , and n are independently an integer of from 1 to 4, and R^1 , R^2 , R^3 and R^4 may be the same or different from the others when k , l , m , and n are an integer of from 2 to 4.

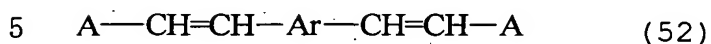
- 5 Specific examples of the biphenylamine compound having the formula (50) include 4'-methoxy-N,N-diphenyl-[1,1'-biphenyl]-4-amine, 4'-methyl-N,N-bis(4-methylphenyl)-[1,1'-biphenyl]-4-amine, 4'-methoxy-N,N-bis(4-methylphenyl)-[1,1'-biphenyl]-4-amine, 10 N,N-bis(3,4-dimethylphenyl)-[1,1'-biphenyl]-4-amine, etc.



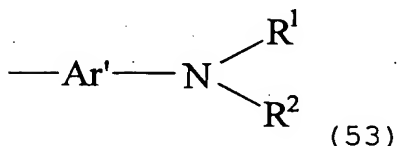
- wherein Ar represents a condensation polycyclic hydrocarbon group having 18 or less carbon atoms which can have a substituent; and R^1 and R^2 independently represent a hydrogen atom, a halogen atom, a substituted or unsubstituted alkyl group, an alkoxy group, or a substituted or unsubstituted phenyl group and n is 1 or 2.

- Specific examples of the triarylamine compound having the formula (51) include N,N-diphenyl-pyrene-1-amine, N,N-di-p-tolyl-pyrene-1-amine, N,N-di-p-tolyl-1-naphthylamine, N,N-di(p-tolyl)-1-phenanthrylamine, 9,9-dimethyl-2-(di-p-tolylamino)fluorene,

N,N,N',N' -tetrakis(4-methylphenyl)-phenanthrene-9,10-diamine,
 N,N,N',N' -tetrakis(3-methylphenyl)-m-phenylenediamine,
 etc.

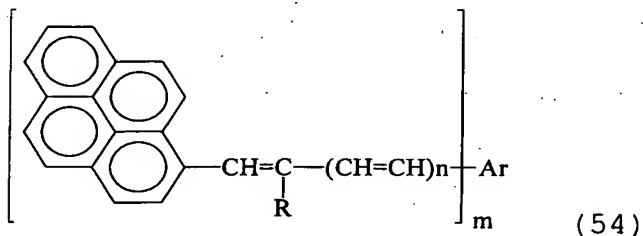


wherein Ar represents a substituted or unsubstituted aromatic hydrocarbon group; and A represents the following formula (53):



10 wherein Ar' represents a substituted or unsubstituted aromatic hydrocarbon group; and R¹ and R² represent substituted or unsubstituted alkyl group or a substituted or unsubstituted aryl group.

Specific examples of the diolefin aromatic compound having
 15 the formula (52) include 1,4-bis(4-diphenylaminostyryl)benzene,
 1,,4-bis [4-di(p-tolyl)aminostyryl] benzene, etc.

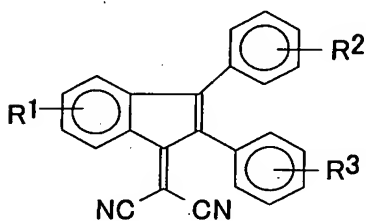


wherein Ar represents a substituted or unsubstituted aromatic
 20 hydrocarbon group; R represents a hydrogen atom, a substituted or unsubstituted alkyl group or a substituted or unsubstituted aryl group; n is 0 or 1; m is 1 or 2; and Ar and R may form a

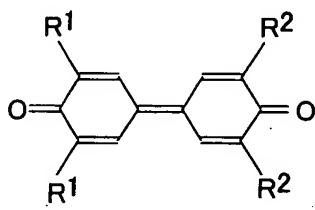
ring when n is 0 and m is 1.

Specific examples of the styrylpyrene compound having the formula (54) include 1-(4-diphenylaminostyryl)pyrene, 1-[4-di(p-tolyl)aminostyryl] pyrene, etc.

5 Specific examples of an electron transport materials include chloranil, bromoanil, tetracyanoethylene, tetracyanoquinodimethane, 2,4,7-trinitro-9-fluorenone, 2,4,5,7-tetranitro-9-fluorenone, 2,4,5,7-tetranitroxanthone, 2,4,8-trinitrothioxanthone, 2,6,8-trinitro-indeno [1,2-b]
10 thiophene-4-one, and 1,3,7-trinitrodibenzothiophene-5,5-dioxide, etc. In addition, electron transport materials having the following formulae (55), (56) and (57) are preferably used.

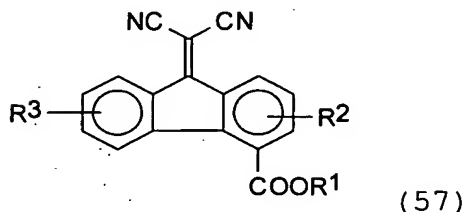


wherein R¹, R² and R³ independently represent a hydrogen atom, a halogen atom, a substituted or unsubstituted alkyl group, an alkoxy group or a substituted or unsubstituted phenyl group;



wherein R¹ and R² independently represent a hydrogen atom, a

substituted or unsubstituted alkyl group; or a substituted or unsubstituted phenyl group;

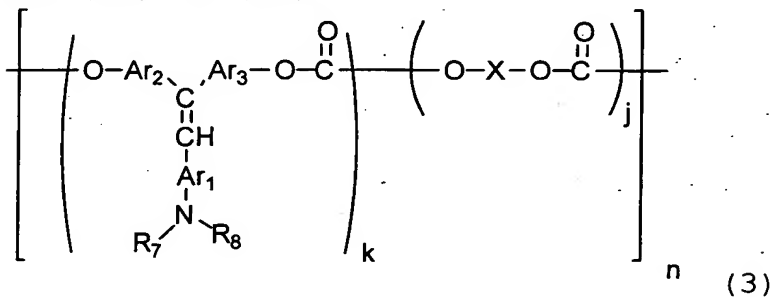


5 wherein R^1 , R^2 and R^3 independently represent a hydrogen atom, a halogen atom, a substituted or unsubstituted alkyl group, an alkoxy group or a substituted or unsubstituted phenyl group.

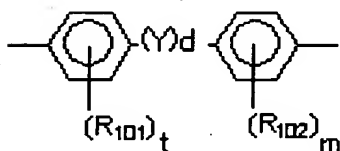
These CTMs can be used alone or in combination.

In the present invention, among the above-mentioned
 10 low-molecular-weight CTMs, the low-molecular-weight CTMs having formulae (1) and (2) are preferably used because of particularly having good transportability and good properties of receiving charges from CGMs. Therefore, an electrophotographic photoreceptor including the
 15 low-molecular-weight CTMs having formulae (1) and (2) in its photosensitive layer has high sensitivity.

Specific examples of the charge transport polymer material include compounds having the following formulae (3), (7), (8) and (58) to (69):

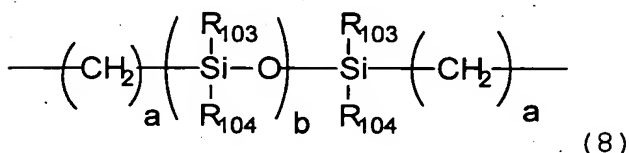


wherein R^7 and R^8 independently represent a substituted or unsubstituted aromatic ring group; Ar^1 , Ar^2 and Ar^3 independently represent an aromatic ring group; k is a number of from 0.1 to 1.0 and j is a number of from 0 to 0.9; n represents a repeating number and is an integer of from 5 to 5,000; and X represents a divalent aliphatic group, a divalent alicyclic group or a divalent group having the following formula (7):



(7)

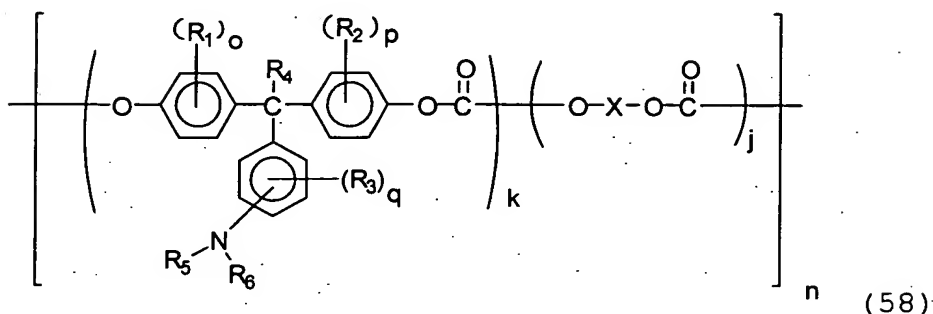
wherein, R^{101} and R^{102} independently represent a substituted or unsubstituted alkyl group, a substituted or unsubstituted aryl group, or a halogen atom; t and m independently represent 0 or an integer of from 1 to 4; d is 0 or 1; and Y represents a linear alkylene group, a branched alkylene group, a cyclic alkylene group, $-O-$, $-S-$, $-SO-$, $-SO_2-$, $-CO-$, $-CO-O-Z-O-CO-$ (Z represents a divalent aliphatic group), or a group having the following formula (8):



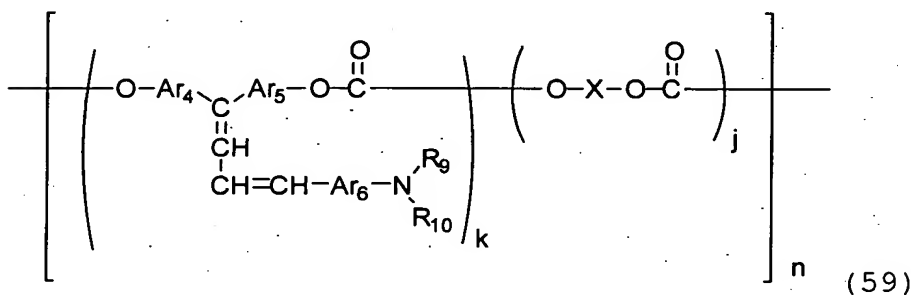
(8)

wherein, a is an integer of from 1 to 20; b is an integer of from 1 to 2,000; and R^{103} and R^{104} independently represent a substituted or unsubstituted alkyl group, or a substituted or

unsubstituted aryl group, and wherein R^{101} , R^{102} , R^{103} and R^{104} may be the same or different from the others.

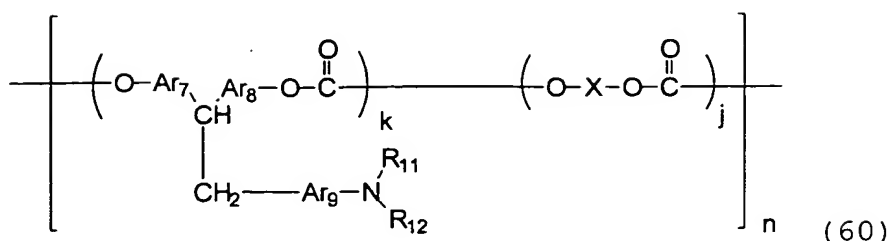


- 5 wherein, R_1 , R_2 and R_3 independently represent a substituted or unsubstituted alkyl group, or a halogen atom; R_4 represents a hydrogen atom, or a substituted or unsubstituted alkyl group; R_5 , and R_6 independently represent a substituted or unsubstituted aryl group; o , p and q independently represent 0 or an integer
10 of from 1 to 4; and X , k , j and n are same in formula (3);

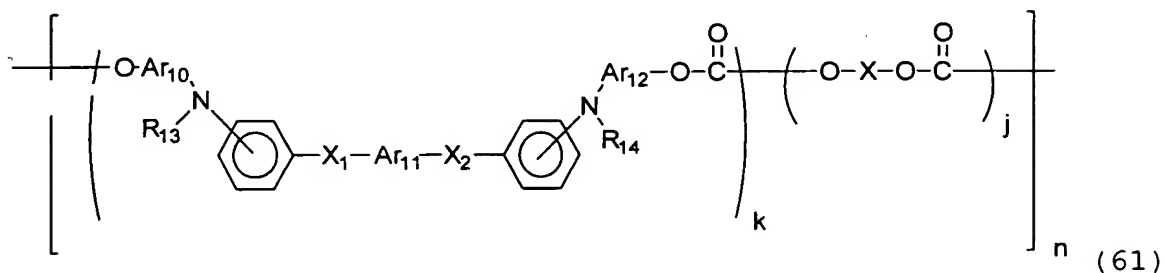


- wherein, R_9 and R_{10} represent a substituted or unsubstituted aryl group; Ar_4 , Ar_5 and Ar_6 independently represent an arylene group;
15 and X , k , j and n are same in formula (3);

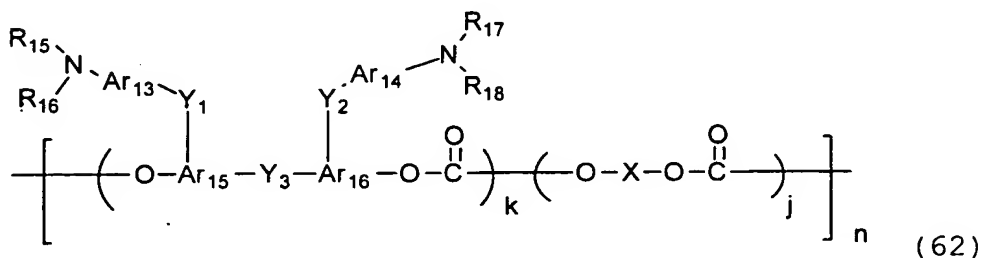
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wherein, R_{11} and R_{12} represent a substituted or unsubstituted aryl group; Ar_7 , Ar_8 and Ar_9 independently represent an arylene group; p is an integer of from 1 to 5; and X , k , j and n are same in formula (3);



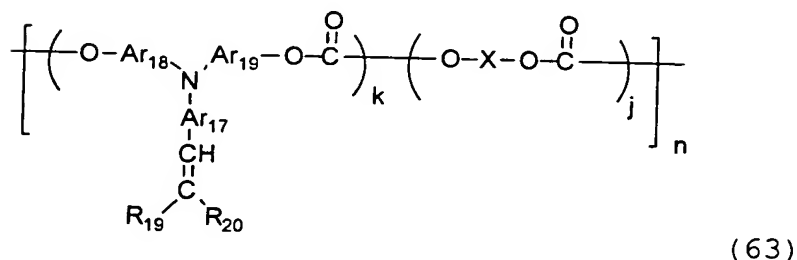
wherein, R_{13} and R_{14} represent a substituted or unsubstituted aryl group; Ar_{10} , Ar_{11} and Ar_{12} independently represent an arylene group; X_1 and X_2 represent a substituted or unsubstituted ethylene group, or a substituted or unsubstituted vinylene group; and X , k , j and n are same in formula (3);



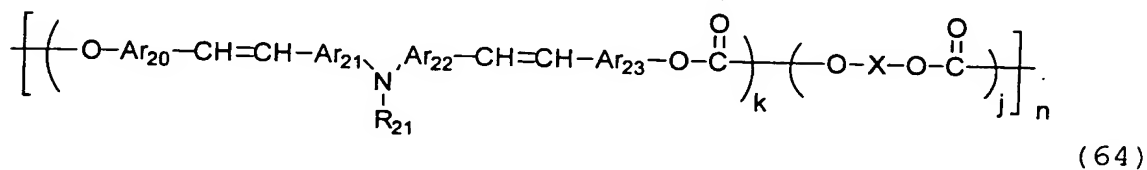
wherein, R_{15} , R_{16} , R_{17} and R_{18} represent a substituted or unsubstituted aryl group; Ar_{13} , Ar_{14} , Ar_{15} and Ar_{16} independently represent an arylene group; Y_1 , Y_2 and Y_3 independently represent

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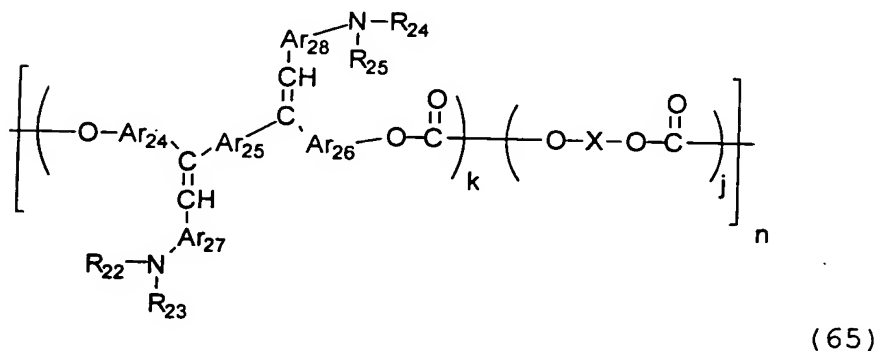
a direct bonding, a substituted or unsubstituted alkylene group, a substituted or unsubstituted cycloalkylene group, a substituted or unsubstituted alkylene ether group, an oxygen atom, a sulfur atom, or a vinylene group; and X, k, j and n are same in formula (3);



wherein, R₁₉ and R₂₀ represent a hydrogen atom, or substituted or unsubstituted aryl group, and R₁₉ and R₂₀ may form a ring; Ar₁₇, Ar₁₈ and Ar₁₉ independently represent an arylene group; and X, k, j and n are same in formula (3);



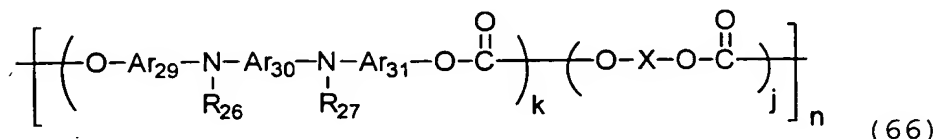
wherein, R₂₁ represents a substituted or unsubstituted aryl group; Ar₂₀, Ar₂₁, Ar₂₂ and Ar₂₃ independently represent an arylene group; and X, k, j and n are same in formula (3);



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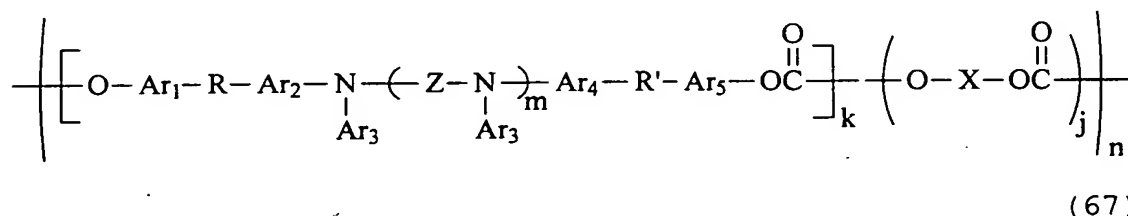
wherein, R_{22} , R_{23} , R_{24} and R_{25} represent a substituted or unsubstituted aryl group; Ar_{24} , Ar_{25} , Ar_{26} , Ar_{27} and Ar_{28} independently represent an arylene group; and X , k , j and n are same in formula (3);

5



wherein, R_{26} and R_{27} independently represent a substituted or unsubstituted aryl group; Ar_{29} , Ar_{30} and Ar_{31} independently represent an arylene group; and X , k , j and n are same in formula

10 (3);



wherein Ar_1 , Ar_2 , Ar_3 , Ar_4 and Ar_5 represent a substituted or unsubstituted aromatic ring group; Z represents an aromatic ring group or $-Ar_6-Za-Ar_6-$; Ar_6 represents a substituted or unsubstituted aromatic ring group; Za represents O , S or an alkylene group; R and R' represent a linear alkylene group or a branched alkylene group; m is 0 or 1; and X , k , j and n are same in formula (3).

In the present invention, among the above-mentioned charge transport polymer materials, the charge transport polymer material having formula (3) is preferably used because of particularly having good abrasion resistance and good

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transportability. Therefore, an electrophotographic photoreceptor including the charge transport polymer material having formula (3) in its photosensitive layer has high durability and sensitivity.

5 Next, layer compositions of the electrophotographic photoreceptor of the present invention will be explained, referring to Figs. 1 to 5.

Fig. 1 is a schematic view illustrating a cross section of a surface of an embodiment of the photoreceptor of the present invention, in which a photosensitive layer 33 including a CGM
10 as the main components is formed on an electroconductive substrate 31.

In Fig. 2, a CGL 35 including a CGM as the main component overlies a CTL 37 including a CTM as the main component on an
15 electroconductive substrate 31.

In Fig. 3, a photosensitive layer 33 including a CGM and a CTM as the main components is formed on an electroconductive substrate 31, and further a protection layer 39 is formed on a surface of the photosensitive layer. In this case, the
20 protection layer 39 may include an amine compound of the present invention.

In Fig. 4, a CGL 35 including a CGM as the main component, a CTL 37 including a CTM as the main component overlying the CGL, and further a protection layer 39 overlying the CTL are
25 formed on an electroconductive substrate 31. In this case, the protection layer 39 may include an amine compound of the present invention.

In Fig. 5, a CTL 37 including a CTM as the main component, a CGL 35 including a CGM as the main component overlying the CTL, and further a protection layer 39 overlying the CGL are formed on an electroconductive substrate 31. In this case, the protection layer 39 may include an amine compound of the present invention.

Suitable materials for use as the electroconductive substrate 31 include materials having a volume resistance not greater than $10^{10} \Omega \cdot \text{cm}$. Specific examples of such materials include plastic cylinders, plastic films or paper sheets, on the surface of which a metal such as aluminum, nickel, chromium, nichrome, copper, gold, silver, platinum and the like, or a metal oxide such as tin oxides, indium oxides and the like, is deposited or sputtered. In addition, a plate of a metal such as aluminum, aluminum alloys, nickel and stainless steel and a metal cylinder, which is prepared by tubing a metal such as the metals mentioned above by a method such as impact ironing or direct ironing, and then treating the surface of the tube by cutting, super finishing, polishing and the like treatments, can be also used as the substrate. Further, endless belts of a metal such as nickel and stainless steel, which have been disclosed in Japanese Laid-Open Patent Publication No. 52-36016, can be also used as the electroconductive substrate 31.

Furthermore, substrates, in which a coating liquid including a binder resin and an electroconductive powder is coated on the supporters mentioned above, can be used as the substrate 31. Specific examples of such an electroconductive

powder include carbon black, acetylene black, powders of metals such as aluminum, nickel, iron, Nichrome, copper, zinc, silver and the like, and metal oxides such as electroconductive tin oxides, ITO and the like. Specific examples of the binder resin
5 include known thermoplastic resins, thermosetting resins and photo-crosslinking resins, such as polystyrene, styrene-acrylonitrile copolymers, styrene-butadiene copolymers, styrene-maleic anhydride copolymers, polyesters, polyvinyl chloride, vinyl chloride-vinyl acetate copolymers,
10 polyvinyl acetate, polyvinylidene chloride, polyarylates, phenoxy resins, polycarbonates, cellulose acetate resins, ethyl cellulose resins, polyvinyl butyral resins, polyvinyl formal resins, polyvinyl toluene, poly-N-vinyl carbazole, acrylic resins, silicone resins, epoxy resins, melamine resins, urethane
15 resins, phenolic resins, alkyd resins and the like resins. Such an electroconductive layer can be formed by coating a coating liquid in which an electroconductive powder and a binder resin are dispersed in a solvent such as tetrahydrofuran, dichloromethane, methyl ethyl ketone, toluene and the like
20 solvent, and then drying the coated liquid.

In addition, substrates, in which an electroconductive resin film is formed on a surface of a cylindrical substrate using a heat-shrinkable resin tube which is made of a combination of a resins such as polyvinyl chloride, polypropylene, polyesters,
25 polyvinylidene chloride, polyethylene, chlorinated rubber and fluorine-containing resins, with an electroconductive material, can be also used as the substrate 31.

Next, the photosensitive layer of the present invention will be explained. In the present invention, the photosensitive layer may be single-layered or a multi-layered. At first, the multi-layered photosensitive layer including the CGL 35 and the CTL 37 will be explained for explanation convenience.

The CGL 35 is a layer including a CGM as the main component. Known CGMs can be used in the CGL 35. Specific examples of the CGM include azo pigments such as CI Pigment Blue 25 (color index CI 21180), CI Pigment Red 41 (CI 21200), CI Acid Red 52 (CI 45100), CI Basic Red 3 (CI 45210), an azo pigment having a carbazole skeleton disclosed in Japanese Laid-Open Patent Publication (JLPP) No. 53-95033, an azo pigment having a distyrylbenzene skeleton disclosed in JLPP No. 53-133445, an azo pigment having a triphenylamine skeleton disclosed in JLPP No. 53-132347, an azo pigment having a dibenzothiophene skeleton disclosed in JLPP No. 54-21728, an azo pigment having an oxadiazole skeleton disclosed in JLPP No. 54-12742, an azo pigment having a fluorenone skeleton disclosed in JLPP No. 54-22834, an azo pigment having a bisstilbene skeleton disclosed in JLPP No. 54-17733, an azo pigment having a distyryloxadiazole skeleton disclosed in JLPP No. 54-2129, an azo pigment having a distyrylcarbazole skeleton disclosed in JLPP No. 54-14967 and an azo pigment having a benzanthrone skeleton; phthalocyanine pigments such as CI Pigment Blue 16 (CI 74100), Y-type oxotitaniumphthalocyanine disclosed in JLPP No. 64-17066, A(β)-type oxotitaniumphthalocyanine, B(α)-type -type oxotitaniumphthalocyanine, I-type oxotitaniumphthalocyanine

disclosed in JLPP No. 11-21466, II-type chlorogalliumphthalocyanine disclosed by Mr. Iijima and others in the 67th spring edition 1B4, 04 published by Chemical Society of Japan in 1994, V-type hydroxygalliumphthalocyanine disclosed Mr. Daimon and others in the 67th spring edition 1B4, 05 published by Chemical Society of Japan in 1994 and X-type metal-free phthalocyanine disclosed in US Patent No. 3,816,118; indigo pigments such as CI Vat Brown 5 (CI 73410) and CI Vat Dye (CI 73030); and perylene pigments such as Algo Scarlet B from Bayer AG and Indanthrene Scarlet R from Bayer AG. These materials can be used alone or in combination.

The CGL 35 can be prepared by dispersing a CGM in a proper solvent optionally together with a binder resin using a ball mill, an attritor, a sand mill or a supersonic dispersing machine, coating the coating liquid on an electroconductive substrate and then drying the coated liquid.

Specific example of the binder resins optionally used in the CGL 35, include polyamides, polyurethanes, epoxy resins, polyketones, polycarbonates, silicone resins, acrylic resins, polyvinyl butyral, polyvinyl formal, polyvinyl ketones, polystyrene, polysulfone, poly-N-vinylcarbazole, polyacrylamide, polyvinyl benzal, polyesters, phenoxy resins, vinyl chloride-vinyl acetate copolymers, polyvinyl acetate, polyphenylene oxide, polyamides, polyvinyl pyridine, cellulose resins, casein, polyvinyl alcohol, polyvinyl pyrrolidone, and the like resins. The content of the binder resin in the CGL 35 is preferably from 0 to 500 parts by weight, and preferably

from 10 to 300 parts by weight, per 100 parts by weight of the CGM. The binder resin can be included either before or after dispersion of the CGM in the solvent.

Specific examples of the solvent include isopropanol, acetone, methyl ethyl ketone, cyclohexanone, tetrahydrofuran, dioxane, ethyl cellosolve, ethyl acetate, methyl acetate, dichloromethane, dichloroethane, monochlorobenzene, cyclohexane, toluene, xylene, ligroin, and the like solvents. In particular, ketone type solvents, ester type solvents and ether type solvents are preferably used. These can be used alone or in combination.

The CGL 35 includes a CGM, a solvent and a binder resin as the main components. Any additives such as a sensitizer, a disperser, a detergent and a silicone oil can be included therein.

The coating liquid can be coated by a coating method such as dip coating, spray coating, bead coating, nozzle coating, spinner coating and ring coating. The CGL 35 preferably has a thickness of from 0.01 to 5 μm , and more preferably from 0.1 to 2 μm .

The CTL 37 is a layer including a CTM as the main component and a compound having an alkylamino group. The CTMs having the above-mentioned formulae (1) to (8) and (36) to (67) are preferably used, and the compound having an alkylamino group CTMs having the above-mentioned formulae (9) to (35) (specifically the above-mentioned compounds Nos. 1 to 34) are preferably used. The CTL 37 is formed by dissolving the CTM,

compound having an alkylamino group and optionally a binder resin in a proper solvent to prepare a coating liquid, coating the coating liquid on the CGL 35 and drying the coating liquid.

Specific examples of the binder resin include

5 thermoplastic resins, thermosetting resins such as polystyrene, styrene-acrylonitrile copolymers, styrene-butadiene copolymers, styrene-maleic anhydride copolymers, polyesters, polyvinyl chloride, vinyl chloride-vinyl acetate copolymers, polyvinyl acetate, polyvinylidene chloride, polyarylates,
10 phenoxy resins, polycarbonates, cellulose acetate resins, ethyl cellulose resins, polyvinyl butyral resins, polyvinyl formal resins, polyvinyl toluene, poly-N-vinyl carbazole, acrylic resins, silicone resins, epoxy resins, melamine resins, urethane resins, phenolic resins, alkyd resins and the like.

15 When a photosensitive layer is formed of the CGL 35 and CTL 37, and the CTM and compound having an alkylamino group are included in the CTL 37, a total content of the CTM and compound having an alkylamino group is preferably from 20 to 300 parts by weight, and more preferably from 40 to 150 parts by weight
20 per 100 parts by weight of the binder resin. The CTL preferably has a thickness not greater than 25 μm in view of resolution of the resultant images and response. The lower limit of the thickness is preferably not less than 5 μm , although it depends on the image forming system (particularly on a charged potential
25 of the electrophotographic photoreceptor).

In addition, the content of the compound having an alkylamino group is preferably from 0.01 to 150 % by weight based

on total weight of the CTM. When less than 0.01 % by weight, the durability against the oxidizing gas of the resultant photoreceptor deteriorates. When greater than 150 % by weight, the residual potential thereof increases.

5 In the present invention, an oxidation potential (Eox1) of the compound having an alkylamino group and that (Eox2) of the CTM satisfy the following relationship (I):

$$\text{Eox1} - \text{Eox2} \geq -0.2 \quad (\text{I})$$

To satisfy the relationship, a compound having an
10 alkylamino group and an oxidation potential (Eox1) which is not far from a fixed oxidation potential (Eox2) of the main CTM is preferably selected. When a threshold, i.e. Eox1 - Eox2, is less than -0.2, the compound having an alkylamino group noticeably becomes a trap against the charge transport (hole)
15 and a bright section potential of the resultant electrophotographic photoreceptor becomes large, and therefore the resultant images do not have a contrast.

Specific examples of a solvent for use in forming the CTL
37 include tetrahydrofuran, dioxane, toluene, dichloromethane,
20 monochlorobenzene, dichloroethane, cyclohexanone, methylethyl ketone, acetone and the like solvents. The CTM can be used alone or in combination in the solvent.

As an antioxidant is preferably included in the CTL 37 and conventional antioxidants mentioned later can be used, and
25 (c) hydroquinone compounds and (f) hindered amine compounds are effectively used in particular.

However, the antioxidant for use in the CTL has a

different purpose from the after-mentioned purpose, and are used to prevent quality alteration of the amine compound of the present invention. Therefore, the antioxidant is preferably included in a CTL coating liquid before the amine compound of the present invention is included therein. The content of the antioxidant is from 0.1 to 200 % by weight based on total weight of the amine compound.

The CTL 37 preferably includes a polymer CTM, which has both a binder resin function and a charge transport function, because the resultant CTL 37 has good abrasion resistance. Suitable charge transport polymer materials include known materials. Among these materials, polycarbonate resins having a triarylamine structure in their main chain and/or side chain are preferably used.

The CTL 37 can be formed by coating a coating liquid in which the CTM alone or the CTM and a binder resin are dissolved or dispersed in a proper solvent on the CGL, and drying the liquid. In addition, the CTL may optionally include two or more of additives such as plasticizers, leveling agents and antioxidants.

As a method of coating the thus prepared coating liquid, a conventional coating method such as a dip coating method, a spray coating method, a bead coating method, a nozzle coating method, a spinner coating method and a ring coating method can be used.

Next, a single-layered photosensitive layer will be explained. A photoreceptor in which the above-mentioned CGM

is dispersed in the binder resin can be used. The photosensitive layer can be formed by coating a coating liquid in which a CGM, a CTM and a binder resin are dissolved or dispersed in a proper solvent, and then drying the coated liquid. In addition, the photosensitive layer may optionally include additives such as plasticizers, leveling agents and antioxidants.

Suitable binder resins include the resins mentioned above in the CTL 37. The resins mentioned above in the CGL 35 can be added as a binder resin. In addition, the polymer CTLs mentioned above can be also used as a binder resin preferably. A content of the CGM is preferably from 5 to 40 parts by weight per 100 parts by weight of the binder resin. A total content of the CGM and the compound having an alkylamino group is preferably from 10 to 45 parts by weight, and more preferably from 20 to 30 parts by weight. Further, the compound having an alkylamino group preferably has a content of from 5 to 100 % by weight per 100 % by weight of the CTM. When less than 5 % by weight, the resultant electrophotographic photoreceptor does not have sufficient resistance against the oxidizing gas. When greater than 100 % by weight, the residual potential of the resultant electrophotographic photoreceptor due to repeated use increases.

A method of satisfying the above-mentioned relationship (I) between an oxidation potential (E_{ox1}) of the compound having an alkylamino group and that (E_{ox2}) of the CTM when the photosensitive layer is single-layered is the same as that for

the above-mentioned photosensitive layer formed of the CGL 35 and CTL 37.

The single-layered photosensitive layer can be formed by coating a coating liquid in which a CGM, a binder resin and
5 a CTM are dissolved or dispersed in a solvent such as tetrahydrofuran, dioxane, dichloroethane, cyclohexane, etc. by a coating method such as a dip coating method, spray coating method, a bead coating method and a ring coating method. The thickness of the photosensitive layer is preferably from 5 to
10 25 μm .

In the photoreceptor of the present invention, an undercoat layer may be formed between the substrate 31 and the photosensitive layer. The undercoat layer includes a resin as a main component. Since a photosensitive layer is typically
15 formed on the undercoat layer by coating a liquid including an organic solvent, the resin in the undercoat layer preferably has good resistance against general organic solvents. Specific examples of such resins include water-soluble resins such as polyvinyl alcohol resins, casein and polyacrylic acid
20 sodium salts; alcohol soluble resins such as nylon copolymers and methoxymethylated nylon resins; and thermosetting resins capable of forming a three-dimensional network such as polyurethane resins, melamine resins, alkyd-melamine resins, epoxy resins and the like. The undercoat layer may include a
25 fine powder of metal oxides such as titanium oxide, silica, alumina, zirconium oxide, tin oxide and indium oxide to prevent occurrence of moiré in the recorded images and to decrease

residual potential of the photoreceptor.

The undercoat layer can be formed by coating a coating liquid using a proper solvent and a proper coating method similarly to those for use in formation of the photosensitive layer mentioned above. The undercoat layer may be formed using a silane coupling agent, titanium coupling agent or a chromium coupling agent. In addition, a layer of aluminum oxide which is formed by an anodic oxidation method and a layer of an organic compound such as polyparaxylylene (parylene) or an inorganic compound such as SiO_2 , SnO_2 , TiO_2 , ITO or CeO_2 which is formed by a vacuum evaporation method is also preferably used as the undercoat layer. The thickness of the undercoat layer is preferably 0 to 5 μm .

In the photoreceptor of the present invention, the protection layer 39 is optionally formed overlying the photosensitive layer. Suitable materials for use in the protection layer 39 include organic compounds having an acid value of from 10 to 400 mgKOH/g such as ABS resins, ACS resins, olefin-vinyl monomer copolymers, chlorinated polyethers, aryl resins, phenolic resins, polyacetal, polyamides, polyester resins, polyamideimide, polyacrylates, polyarylsulfone, polybutylene, polybutylene terephthalate, polycarbonate, polyethersulfone, polyethylene, polyethylene terephthalate, polyimides, acrylic resins, polymethylpentene, polypropylene, polyphenyleneoxide, polysulfone, polystyrene, AS resins, butadiene-styrene copolymers, polyurethane, polyvinyl chloride, polyvinylidene chloride, epoxy resins and the like, because of

preventing an increase of residual potential of the resultant photoreceptor. Among these materials, the polycarbonate resin and the polyarylate resin are preferably and effectively used in terms of dispersibility of a filler, decrease of residual potential and coating defect of the resultant photoreceptor.

The protection layer 39 preferably includes a filler for the purpose of improving abrasion resistance thereof. Suitable fillers include highly-insulative fillers having a pH not less than 5 and a dielectric constant not less than 5 such as titanium oxide, alumina, zinc oxide and zirconium oxide.

The protection layer 39 can be formed by dispersing a binder resin, a filler material, etc. in a proper solvent with a ball mill, an attritor, a sand mill or an ultrasonic to prepare a dispersion liquid; coating the dispersion liquid on a photosensitive layer; and drying the dispersion liquid.

As a solvent for use in forming the protection layer, tetrahydrofuran, dioxane, toluene, dichloromethane, monochlorobenzene, dichloroethane, cyclohexanone, methyl ethyl ketone, acetone and the like solvents which are all used in the CTL 37 can be used. However, a high-viscosity solvent is preferably used in dispersion, and a high-volatile solvent is preferably used in coating. When such a solvent as satisfies the conditions is not available, a mixture of two or more of solvents having each property can be used, which occasionally improves dispersibility of the filler and decreases residual potential of the resultant photoreceptor.

Further, the protection layer 39 may include the compound

having an alkylamino group of the present invention. A CTM and the compound having an alkylamino group are preferably and effectively included therein to decrease residual potential of the resultant photoreceptor and to improve quality of the resultant images.

As a method of forming the protection layer, a conventional coating method such as a dip coating method, a spray coating method, a bead coating method, a nozzle coating method, a spinner coating method and ring coating method can be used. In particular, the spray coating method is preferably used in terms of coated film uniformity.

In the photoreceptor of the present invention, an intermediate layer may be formed between the photosensitive layer and the protection layer. The intermediate layer includes a resin as a main component. Specific examples of the resin include polyamides, alcohol soluble nylons, water-soluble polyvinyl butyral, polyvinyl butyral, polyvinyl alcohol, and the like. The intermediate layer can be formed by one of the above-mentioned known coating methods. The thickness of the intermediate layer is preferably from 0.05 to 2 μm .

In the photoreceptor of the present invention, antioxidants, plasticizers, lubricants, ultraviolet absorbents and leveling agents can be included in each layer such as the CGL, CTL, undercoat layer, protection layer and intermediate layer for environmental improvement, above all for the purpose of preventing decrease of photosensitivity and increase of residual potential. Such compounds will be shown

as follows.

Suitable antioxidants for use in the layers of the photoreceptor include the following compounds but are not limited thereto.

5 (a) Phenolic compounds

2,6-di-t-butyl-p-cresol, butylated hydroxyanisole,
2,6-di-t-butyl-4-ethylphenol,
n-octadecyl-3-(4'-hydroxy-3',5'-di-t-butylphenol),
2,2'-methylene-bis-(4-methyl-6-t-butylphenol),
10 2,2'-methylene-bis-(4-ethyl-6-t-butylphenol),
4,4'-thiobis-(3-methyl-6-t-butylphenol),
4,4'-butylidenebis-(3-methyl-6-t-butylphenol),
1,1,3-tris-(2-methyl-4-hydroxy-5-t-butylphenyl)butane,
1,3,5-trimethyl-2,4,6-tris(3,5-di-t-butyl-4-hydroxybenzyl)b
15 enzene,
tetrakis-[methylene-3-(3',5'-di-t-butyl-4'-hydroxyphenyl)pr
opionate]methane,
bis[3,3'-bis(4'-hydroxy-3'-t-butylphenyl)butyric
acid]glycol ester, tocophenol compounds, and the like.

20 (b) Paraphenylenediamine compounds

N-phenyl-N'-isopropyl-p-phenylenediamine,
N,N'-di-sec-butyl-p-phenylenediamine,
N-phenyl-N-sec-butyl-p-phenylenediamine,
N,N'-di-isopropyl-p-phenylenediamine,
25 N,N'-dimethyl-N,N'-di-t-butyl-p-phenylenediamine, and the
like.

(c) Hydroquinone compounds

2,5-di-t-octylhydroquinone, 2,6-didodecylhydroquinone,
2-dodecylhydroquinone, 2-dodecyl-5-chlorohydroquinone, 2-t-
octyl-5-methylhydroquinone,
2-(2-octadecenyl)-5-methylhydroquinone and the like.

5 (d) Organic sulfur-containing compounds

Dilauryl-3,3'-thiodipropionate,
distearyl-3,3'-thiodipropionate,
ditetradecyl-3,3'-thiodipropionate, and the like.

(e) Organic phosphorus-containing compounds

10 Triphenylphosphine, tri(nonylphenyl)phosphine,
tri(dinonylphenyl)phosphine, tricresylphosphine,
tri(2,4-dibutylphenoxy)phosphine and the like.

Suitable plasticizers for use in the layers of the
photoreceptor include the following compounds but are not limited
15 thereto:

(a) Phosphoric acid esters plasticizers

Triphenyl phosphate, tricresyl phosphate, trioctyl
phosphate, octyldiphenyl phosphate, trichloroethyl phosphate,
cresyldiphenyl phosphate, tributyl phosphate, tri-2-ethylhexyl
20 phosphate, triphenyl phosphate, and the like.

(b) Phthalic acid esters plasticizers

Dimethyl phthalate, diethyl phthalate, diisobutyl
phthalate, dibutyl phthalate, diheptyl phthalate,
di-2-ethylhexyl phthalate, diisooctyl phthalate, di-n-octyl
25 phthalate, dinonyl phthalate, diisononyl phthalate, diisodecyl
phthalate, diundecyl phthalate, ditridecyl phthalate,
dicyclohexyl phthalate, butylbenzyl phthalate, butyllauryl

phthalate, methyloleyl phthalate, octyldecyl phthalate, dibutyl fumarate, dioctyl fumarate, and the like.

(c) Aromatic carboxylic acid esters plasticizers

5 Trioctyl trimellitate, tri-n-octyl trimellitate, octyl oxybenzoate, and the like.

(d) Dibasic fatty acid esters plasticizers

 Dibutyl adipate, di-n-hexyl adipate, di-2-ethylhexyl adipate, di-n-octyl adipate, n-octyl-n-decyl adipate, diisodecyl adipate, dialkyl adipate, dicapryl adipate, 10 di-2-ethylhexyl azelate, dimethyl sebacate, diethyl sebacate, dibutyl sebacate, di-n-octyl sebacate, di-2-ethylhexyl sebacate, di-2-ethoxyethyl sebacate, dioctyl succinate, diisodecyl succinate, dioctyl tetrahydrophthalate, di-n-octyl tetrahydrophthalate, and the like.

15 (e) Fatty acid ester derivatives

 Butyl oleate, glycerin monooleate, methyl acetylricinolate, pentaerythritol esters, dipentaerythritol hexaesters, triacetin, tributyrin, and the like.

(f) Oxyacid esters plasticizers

20 Methyl acetylricinolate, butyl acetylricinolate, butylphthalylbutyl glycolate, tributyl acetylcitrate, and the like.

(g) Epoxy plasticizers

 Epoxydized soybean oil, epoxydized linseed oil, butyl 25 epoxystearate, decyl epoxystearate, octyl epoxystearate, benzyl epoxystearate, dioctylepoxyhexahydrophthalate, didecyl epoxyhexahydrophthalate, and the like.

(h) Dihydric alcohol esters plasticizers

Diethylene glycol dibenzoate, triethylene glycol di-2-ethylbutyrate, and the like.

(i) Chlorine-containing plasticizers

5 Chlorinated paraffin, chlorinated diphenyl, methyl esters of chlorinated fatty acids, methyl esters of methoxychlorinated fatty acids, and the like.

(j) Polyester plasticizers

Polypropylene adipate, polypropylene sebacate,
10 acetylated polyesters, and the like.

(k) Sulfonic acid derivatives

P-toluene sulfonamide, o-toluene sulfonamide, p-toluene sulfoneethylamide, o-toluene sulfoneethylamide, toluene sulfone-N-ethylamide, p-toluene sulfone-N-cyclohexylamide,
15 and the like.

(l) Citric acid derivatives

Triethyl citrate, triethyl acetylcitrate, tributyl citrate, tributyl acetylcitrate, tri-2-ethylhexyl acetylcitrate, n-octyldecyl acetylcitrate, and the like.

20 (m) Other compounds

Terphenyl, partially hydrated terphenyl, camphor, 2-nitro diphenyl, dinonyl naphthalene, methyl abietate, and the like.

Suitable lubricants for use in the layers of the
25 photoreceptor include the following compounds but are not limited thereto.

(a) Hydrocarbon compounds

Liquid paraffins, paraffin waxes, micro waxes, low molecular weight polyethylenes, and the like.

(b) Fatty acid compounds

5 Lauric acid, myristic acid, palmitic acid, stearic acid, arachidic acid, behenic acid, and the like.

(c) Fatty acid amide compounds

Stearic acid amide, palmitic acid amide, oleic acid amide, methylenebisstearamide, ethylenebisstearamide, and the like.

(d) Ester compounds

10 Lower alcohol esters of fatty acids, polyhydric alcohol esters of fatty acids, polyglycol esters of fatty acids, and the like.

(e) Alcohol compounds

15 Cetyl alcohol, stearyl alcohol, ethylene glycol, polyethylene glycol, polyglycerol, and the like.

(f) Metallic soaps

Lead stearate, cadmium stearate, barium stearate, calcium stearate, zinc stearate, magnesium stearate, and the like.

(g) Natural waxes

20 Carnauba wax, candelilla wax, beeswax, spermaceti, insect wax, montan wax, and the like.

(h) Other compounds

Silicone compounds, fluorine compounds, and the like.

25 Suitable ultraviolet absorbing agents for use in the layers of the photoreceptor include the following compounds but are not limited thereto.

(a) Benzophenone compounds

2-hydroxybenzophenone, 2,4-dihydroxybenzophenone,
2,2',4-trihydroxybenzophenone,
2,2',4,4'-tetrahydroxybenzophenone,
2,2'-dihydroxy-4-methoxybenzophenone, and the like.

5 (b) Salicylate compounds

Phenyl salicylate,
2,4-di-t-butylphenyl-3,5-di-t-butyl-4-hydroxybenzoate, and
the like.

(c) Benzotriazole compounds

10 (2'-hydroxyphenyl)benzotriazole,
(2'-hydroxy-5'-methylphenyl)benzotriazole and
(2'-hydroxy-3'-t-butyl-5'-methylphenyl)-5-chlorobenzotriazo
le.

(d) Cyano acrylate compounds

15 Ethyl-2-cyano-3,3-diphenyl acrylate,
methyl-2-carbomethoxy-3-(paramethoxy) acrylate, and the like.

(e) Quenchers (metal complexes)

Nickel(2,2'-thiobis(4-t-octyl)phenolate)-n-butylamine,
nickeldibutyldithiocarbamate,
20 cobaltdicyclohexyldithiophosphate, and the like.

(f) HALS (hindered amines)

Bis(2,2,6,6-tetramethyl-4-piperidyl)sebacate,
bis(1,2,2,6,6-pentamethyl-4-piperidyl)sebacate,
1-[2-{3-(3,5-di-t-butyl-4-hydroxyphenyl)propionyloxy} ethyl]
25 -4-{3-(3,5-di-t-butyl-4-hydroxyphenyl)propionyloxy}-2,2,6,6
-tetramethylpyridine,
8-benzyl-7,7,9,9-tetramethyl-3-octyl-1,3,8-triazaspiro[4,5]

undecane-2,4-dione,

4-benzoyloxy-2,2,6,6-tetramethylpiperidine, and the like.

Next, the image forming method and apparatus of the present invention will be explained, referring to drawings.

5 Specifically, the image forming method typified by an electrophotographic image forming method and the image forming apparatus typified by an electrophotographic image forming apparatus will be explained.

Fig. 6 is a schematic view for explaining the
10 electrophotographic method and apparatus of the present invention, and a modified embodiment as mentioned below belongs to the present invention.

In Fig. 6, a photoreceptor 1 includes at least a photosensitive layer and the most surface layer includes a filler.
15 The photoreceptor 1 is drum-shaped, and may be sheet-shaped or endless-belt shaped. Any known chargers such as a corotron, a scorotron, a solid state charger and a charging roller can be used for a charger 3, a pre-transfer charger 7, a transfer charge 10, a separation charger 11 and a pre-cleaning charger
20 13.

The above-mentioned chargers can be used as transfer means, and typically a combination of the transfer charger and the separation charger is effectively used.

Suitable light sources for use in the imagewise light
25 irradiating device 5 and the discharging lamp 2 include fluorescent lamps, tungsten lamps, halogen lamps, mercury lamps, sodium lamps, light emitting diodes (LEDs), laser diodes (LDs),

light sources using electroluminescence (EL) and the like. In addition, in order to obtain light having a desired wave length range, filters such as sharp-cut filters, band pass filters, near-infrared cutting filters, dichroic filters, interference
5 filters, color temperature converting filters and the like can be used.

The above-mentioned light sources can be used for not only the processes mentioned above and illustrated in Fig. 6, but also other processes, such as a transfer process, a discharging
10 process, a cleaning process, a pre-exposure process, which include light irradiation to the photoreceptor.

When the toner image formed on the photoreceptor 1 by a developing unit 6 is transferred onto a transfer sheet 9, all of the toner image are not transferred thereon, and residual
15 toner particles remain on the surface of the photoreceptor 1. The residual toner is removed from the photoreceptor by a fur brush 14 and a blade 15. The residual toner remaining on the photoreceptor 1 can be removed by only a cleaning brush. Suitable
20 cleaning brushes include known cleaning brushes such as fur brushes and mag-fur brushes.

When the photoreceptor which is previously charged positively is exposed to imagewise light, an electrostatic latent image having a positive or negative charge is formed on the photoreceptor. When the latent image having a positive charge
25 is developed with a toner having a negative charge, a positive image can be obtained. In contrast, when the latent image having a positive charge is developed with a toner having a positive

charge, a negative image (i.e., a reversal image) can be obtained.

As the developing method, known developing methods can be used. In addition, as the discharging methods, known discharging methods can be also used.

5 Fig. 7 is a schematic view for explaining another embodiment of the electrophotographic apparatus and method of the present invention. A photoreceptor 21 includes at least a photosensitive layer and the most surface layer includes a filler. The photoreceptor is rotated by rollers 22a and 22b.

10 Charging using a charger 23, imagewise exposure using an imagewise light irradiating device 24, developing using a developing unit (not shown), transferring using a transfer charger 25, pre-cleaning using a light source 26, cleaning using a cleaning brush 27 and discharging using a discharging light
15 source 28 are repeatedly performed. In Fig. 7, the pre-cleaning light irradiating is performed from the side of the substrate of the photoreceptor 21. In this case, the substrate has to be light-transmissive.

The image forming apparatus of the present invention is
20 not limited to the image forming units as shown in Figs. 6 and 7. For example, although the pre-cleaning light irradiation is performed from the substrate side in Fig. 7, the pre-cleaning light irradiating operation can be performed from the photosensitive layer side of the photoreceptor. In addition,
25 the light irradiation in the light image irradiating process and the discharging process may be performed from the substrate side of the photoreceptor

As light irradiation processes, the imagewise irradiation process, pre-cleaning irradiation process, and discharging light irradiation are illustrated. In addition, a pre-transfer light irradiation and a preliminary light irradiation before the imagewise light irradiation, and other known light irradiation processes may also be performed on the photoreceptor.

The above-mentioned image forming unit may be fixedly set in a copier, a facsimile or a printer. However, the image forming unit may be set therein as a process cartridge. The process cartridge means an image forming unit (or device) which includes a photoreceptor, a charger, an imagewise light irradiator, an image developer, an image transferer, a cleaner, and a discharger. Various process cartridges can be used in the present invention. Fig. 8 illustrates an embodiment of the process cartridge.

Having generally described this invention, further understanding can be obtained by reference to certain specific examples which are provided herein for the purpose of illustration only and are not intended to be limiting. In the descriptions in the following examples, the numbers represent weight ratios in parts, unless otherwise specified.

EXAMPLES

Examples 1 to 26 and Comparative Examples 1 to 8

An undercoat coating liquid, a charge generation coating liquid and charge transport coating liquid, which have the

following formulations, were coated in this order on an aluminium cylinder by a dip coating method and dried to prepare photoreceptors 1 to 34 having an undercoat layer of 3.5 μm thick, a CGL of 0.2 μm thick, a CTL of 23 μm thick.

5 Undercoat layer coating liquid

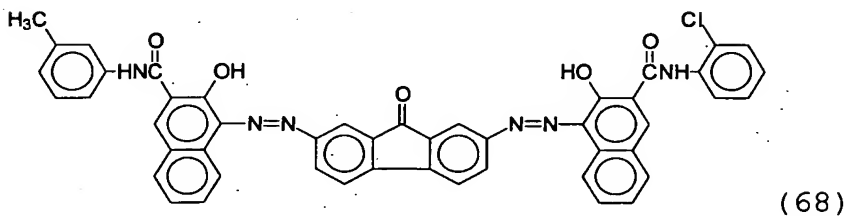
Titanium dioxide powder	400
Melamine resin	65
Alkyd resin	120
2-butanone	400

10

CGL coating liquid

Fluorenone bisazo pigment	12
---------------------------	----

having the following formula (68)



15

Polyvinyl butyral	5
2-butanone	200
Cyclohexanone	400

20 CTL coating liquid

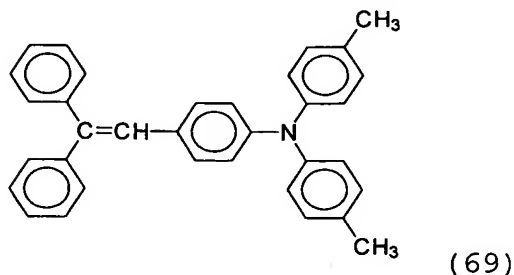
Polycarbonate resin	10
(Z polyca from Teijin Chemicals Ltd.)	
The compounds having an alkylamino group	1
No. 1 to 34	

CTM having the following formula (69)

9

and an oxidation potential of 0.76

(V vs. SCE)



5 Tetrahydrofuran

100

After the thus prepared each photoreceptor 1 to 34 was installed in a process cartridge for electrophotography and the cartridge was installed in a modified copier imagio MF2200 from Ricoh Company, ltd. having a scorotron type corona charger an
 10 imagewise light source of a LD having a wavelength of 655 nm, in which the photoreceptor had a dark portion potential of 800 (-V) and an image surface illuminance of 0.45 ($\mu\text{j}/\text{cm}^2$), a bright portion potential was measured. Further, 100,000 images were continuously produced, and the initial image and the image after
 15 100,000 images were produced were evaluated. The results are shown in Table 1. In addition, a graph in which a difference (ΔE) between an oxidation potential of the compound having an alkylamino group and that of the CTM and the initial bright portion potential (VL) of the photoreceptor are plotted is shown in Fig.

20 9.

Table 1

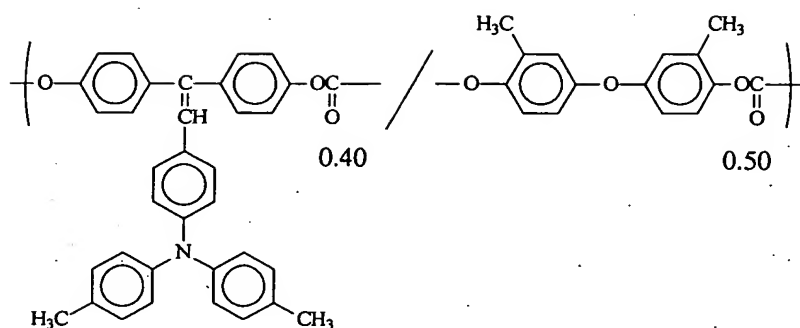
			Initial	After 100,000

	No. of compound	ΔE (V vs. SCE)	VL (-V)	Image quality	Image quality
Com. Ex. 1	1	-0.240	479	Low image density	Low image density
Ex. 1	2	-0.155	133	Good	Good
Com. Ex. 2	3	-0.260	709	Image was not produced	-
Com. Ex. 3	4	-0.320	769	Image was not produced	-
Ex. 2	5	-0.105	117	Good	Good
Com. Ex. 4	6	-0.240	571	Image was not produced	-
Com. Ex. 5	7	-0.210	385	Low image density	Low image density
Ex. 3	8	-0.110	124	Good	Good
Ex. 4	9	-0.120	115	Good	Good
Ex. 5	10	-0.100	113	Good	Good
Ex. 6	11	-0.100	130	Good	Good
Ex. 7	12	-0.135	112	Good	Good
Ex. 8	13	0.115	120	Good	Good
Ex. 9	14	-0.100	102	Good	Good
Ex. 10	15	0.020	105	Good	Good
Ex. 11	16	-0.010	115	Good	Good
Ex. 12	17	-0.160	150	Good	Good
Ex. 13	18	-0.135	131	Good	Good
Com. Ex.	19	-0.365	780	Image was not	-

6				produced	
Ex. 14	20	-0.100	105	Good	Good
Ex. 15	21	-0.140	113	Good	Good
Com. Ex. 7	22	-0.215	318	Good	Low image density
Ex. 16	23	0.000	122	Good	Good
Com. Ex. 8	24	-0.300	785	Image was not produced	-
Ex. 17	25	0.025	101	Good	Good
Ex. 18	26	-0.020	101	Good	Good
Ex. 19	27	-0.010	96	Good	Good
Ex. 20	28	-0.160	109	Good	Good
Ex. 21	29	0.095	76	Good	Good
Ex. 22	30	-0.080	78	Good	Good
Ex. 23	31	-0.140	83	Good	Good
Ex. 24	32	-0.010	102	Good	Good
Ex. 25	33	0.020	95	Good	Good
Ex. 26	34	0.010	105	Good	Good

Examples 27 to 52 and Comparative Examples 9 to 16

5 The procedures for preparation and evaluation of the photoreceptor in Example 1 were repeated to prepare and evaluate photoreceptors 35 to 68 except for changing 9 parts of the CTM and 10 parts of the polycarbonate resin included in the CTL binder resin to 19 parts of a charge transport polymer material having the following formula (70) and an oxidation potential of 0.780 (V. vs. SCE). The results are shown in Table 2 and Fig. 10.



(70)

5

Table 2

	No. of compound	ΔE (V vs. SCE)	Initial		After 100,000
			VL (-V)	Image quality	Image quality
Com. Ex. 9	1	-0.260	519	Low image density	Image was not produced
Ex. 27	2	-0.175	168	Good	Good
Com. Ex. 10	3	-0.260	719	Image was not produced	-
Com. Ex. 11	4	-0.280	779	Image was not produced	-
Ex. 28	5	-0.340	137	Good	Good
Com. Ex. 12	6	-0.125	611	Image was not produced	-
Com. Ex. 13	7	-0.260	285	Good	Low image density

Ex. 29	8	-0.230	134	Good	Good
Ex. 30	9	-0.130	147	Good	Good
Ex. 31	10	-0.140	143	Good	Good
Ex. 32	11	-0.120	160	Good	Good
Ex. 33	12	-0.120	147	Good	Good
Ex. 34	13	-0.155	160	Good	Good
Ex. 35	14	-0.120	102	Good	Good
Ex. 36	15	0.000	125	Good	Good
Ex. 37	16	-0.030	130	Good	Good
Ex. 38	17	-0.180	180	Good	Good
Ex. 39	18	-0.155	161	Good	Good
Com. Ex. 14	19	-0.385	800	Image was not produced	-
Ex. 40	20	-0.120	123	Good	Good
Ex. 41	21	-0.160	137	Good	Good
Com. Ex. 15	22	-0.235	349	Good	Low image density
Ex. 42	23	-0.020	152	Good	Good
Com. Ex. 16	24	-0.320	772	Image was not produced	-
Ex. 43	25	0.005	131	Good	Good
Ex. 44	26	-0.040	131	Good	Good
Ex. 45	27	-0.030	122	Good	Good
Ex. 46	28	-0.180	129	Good	Good
Ex. 47	29	0.075	128	Good	Good
Ex. 48	30	-0.100	106	Good	Good

Ex. 49	31	-0.160	113	Good	Good
Ex. 50	32	-0.030	113	Good	Good
Ex. 51	33	0.000	121	Good	Good
Ex. 52	34	-0.010	135	Good	Good

Examples 53 to 83 and Comparative Examples 17 to 19

The procedures for preparation and evaluation of the photoreceptor in Example 1 were repeated to prepare and evaluate photoreceptors 69 to 102 except for changing the CGL coating liquid and CTL coating liquid to a CGL coating liquid and a CTL coating liquid having the following formulations respectively. The results are shown in Table 3 and Fig. 12.

10 CGL coating liquid

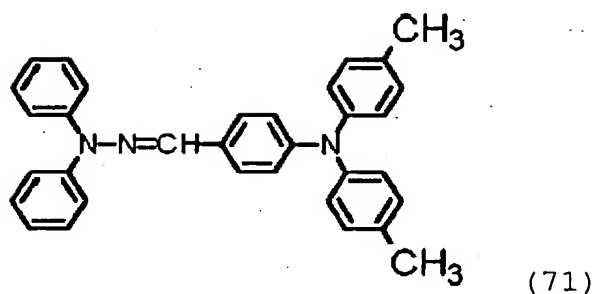
Oxotitaniumphthalocyanine 8
 having the powder XD spectrum in Fig. 11
 Polyvinylbutyral 5
 2-butanone 400

15

CTL coating liquid

Polycarbonate resin 10
 (Z polyca from Teijin Chemicals Ltd.)
 The compounds having an alkylamino group 1
 No. 1 to 34
 CTM having the following formula (71) 7
 and an oxidation potential of 0.675
 (V vs. SCE)

20



Toluene

70

Table 3

	No. of compound	ΔE (V vs. SCE)	Initial		After 100,000
			VL (-V)	Image quality	Image quality
Ex. 53	1	-0.155	94	Good	Good
Ex. 54	2	-0.070	82	Good	Good
Ex. 55	3	-0.175	112	Good	Good
Com. Ex. 17	4	-0.235	423	Low image density	Low image density
Ex. 56	5	-0.020	108	Good	Good
Ex. 57	6	-0.155	129	Good	Good
Ex. 58	7	-0.125	123	Good	Good
Ex. 59	8	-0.025	956	Good	Good
Ex. 60	9	-0.035	116	Good	Good
Ex. 61	10	-0.015	94	Good	Good
Ex. 62	11	-0.015	78	Good	Good
Ex. 63	12	-0.050	65	Good	Good
Ex. 64	13	0.200	105	Good	Good

Ex. 65	14	-0.015	83	Good	Good
Ex. 66	15	0.105	64	Good	Good
Ex. 67	16	0.075	58	Good	Good
Ex. 68	17	-0.075	101	Good	Good
Ex. 69	18	-0.050	73	Good	Good
Com. Ex. 18	19	-0.280	633	Image was not produced	-
Ex. 70	20	-0.015	58	Good	Good
Ex. 71	21	-0.055	87	Good	Good
Ex. 72	22	-0.130	97	Good	Low image density
Ex. 73	23	0.085	81	Good	Good
Com. Ex. 19	24	-0.215	382	Low image density	Low image density
Ex. 74	25	0.110	68	Good	Good
Ex. 75	26	0.065	60	Good	Good
Ex. 76	27	0.075	100	Good	Good
Ex. 77	28	-0.075	95	Good	Good
Ex. 78	29	0.180	110	Good	Good
Ex. 79	30	0.005	97	Good	Good
Ex. 80	31	-0.055	73	Good	Good
Ex. 81	32	0.075	85	Good	Good
Ex. 82	33	0.105	55	Good	Good
Ex. 83	34	0.095	102	Good	Good

Examples 84 to 109 and Comparative Examples 20 to 27

An undercoat coating liquid, a charge generation coating liquid and charge transport coating liquid, which have the following formulations, were coated in this order on an aluminium cylinder by a dip coating method and dried from an undercoat layer of 3.5 μm thick, a CGL of 0.2 μm thick, a CTL of 23 μm thick on the aluminium cylinder.

Undercoat layer coating liquid

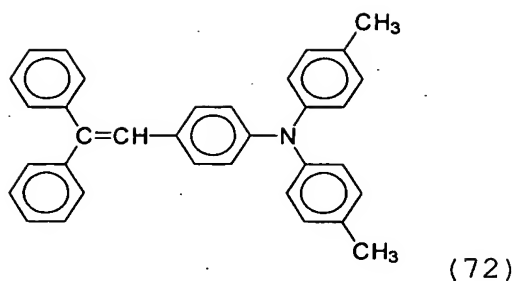
	Titanium dioxide powder	400
10	Melamine resin	65
	Alkyd resin	120
	2-butanone	400

CGL coating liquid

15	Oxotitaniumphthalocyanine	8
	having the powder XD spectrum in Fig. 13	
	Polyvinylbutyral	5
	2-butanone	400

20 CTL coating liquid

	Polycarbonate resin	10
	(Z polyca from Teijin Chemicals Ltd.)	
	CTM having the following formula (72)	9



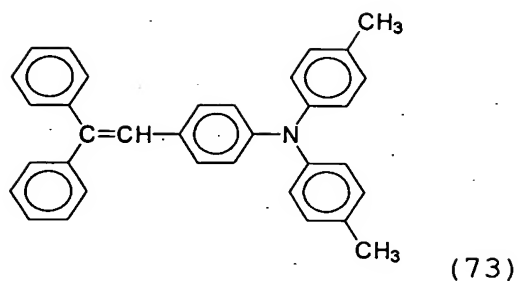
Tetrahydrofuran

100

Further, a protection layer having a thickness of about 4 μm is formed on the CTL by spraying a protection layer coating liquid having the following formulation onto the CTL to prepare photoreceptors 103 to 136. The evaluation results of the photoreceptors are shown in Table 4 and Fig. 14.

Protection layer coating liquid

10	Alumina having an average primary particle diameter of 0.3 μm from Sumitomo Chemical Co., Ltd.	2
	The compounds having an alkylamino group No. 1 to 34	0.5
15	Unsaturated polycarbonate polymer solution having an acid value of 180 mg KOH/g from BYK Chemie GmbH	0.02
	CTM having the following formula (73) and an oxidation potential of 0.76 (V vs. SCE)	3.5
20		



Polycarbonate resin

6

(Z polycarbonate from Teijin Chemicals Ltd.)

Tetrahydrofuran

220

5

Cyclohexanone

80

Table 4

	No. of compound	ΔE (V vs. SCE)	Initial		After 100,000
			VL (-V)	Image quality	Image quality
Com. Ex. 20	1	-0.240	600	Image was not produced	-
Com. Ex. 21	2	-0.155	237	Good	Good
Com. Ex. 22	3	-0.260	720	Image was not produced	-
Com. Ex. 23	4	-0.320	787	Image was not produced	-
Ex. 84	5	-0.105	221	Good	Good
Com. Ex. 24	6	-0.240	682	Image was not produced	-

Ex. 85	7	-0.210	496	Low image density	Low image density
Ex. 86	8	-0.110	220	Good	Good
Ex. 87	9	-0.120	221	Good	Good
Ex. 88	10	-0.100	221	Good	Good
Ex. 89	11	-0.100	222	Good	Good
Ex. 90	12	-0.135	262	Good	Good
Ex. 91	13	0.115	202	Good	Good
Ex. 92	14	-0.100	252	Good	Good
Ex. 93	15	0.020	211	Good	Good
Ex. 94	16	-0.010	218	Good	Good
Ex. 95	17	-0.160	248	Good	Good
Ex. 96	18	-0.135	235	Good	Good
Com. Ex. 25	19	-0.365	765	Image was not produced	-
Ex. 97	20	-0.100	255	Good	Good
Ex. 98	21	-0.140	265	Good	Good
Com. Ex. 26	22	-0.215	448	Good	Low image density
Ex. 99	23	0.000	227	Good	Good
Com. Ex. 27	24	-0.300	789	Image was not produced	-
Ex. 100	25	0.025	216	Good	Good
Ex. 101	26	-0.020	203	Good	Good
Ex. 102	27	-0.010	199	Good	Good
Ex. 103	28	-0.160	216	Good	Good

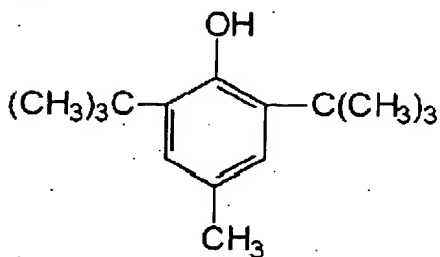
Ex. 104	29	0.095	184	Good	Good
Ex. 105	30	-0.080	184	Good	Good
Ex. 106	31	-0.140	198	Good	Good
Ex. 107	32	-0.010	204	Good	Good
Ex. 108	33	0.020	174	Good	Good
Ex. 109	34	0.010	187	Good	Good

Comparative Example 28

The procedures for preparation and evaluation of the photoreceptor in Example 1 were repeated to prepare and evaluate a comparative photoreceptors 1 except for not adding the compound having an alkylamino group into the CTL coating liquid. The results are shown in Table 5.

Comparative Example 29

The procedures for preparation and evaluation of the photoreceptor in Example 1 were repeated to prepare and evaluate a comparative photoreceptors 2 except for changing the compound having an alkylamino group into a hindered phenol antioxidant having the following formula (74). The results are shown in Table 5.



(74)

Table 5

	Compara- tive Photo- receptor No.	Initial		After 100,000
		VL (-V)	Image quality	Image quality
Com. Ex. 28	1	100	Good	Image resolution lowered (moderate)
Com. Ex. 29	2	545	Low image density	Image density lowered (large) and illegible

The results show that an electrophotographic photoreceptor including an electroconductive substrate and a photosensitive layer on the electroconductive substrate, wherein the photosensitive layer includes at least a compound having a substituted or unsubstituted alkylamino group and a charge transport material, and wherein an oxidation potential (Eox1) of the substituted or unsubstituted alkylamino group and an oxidation potential (Eox2) of the charge transport material satisfy the following relationship (I) has a high sensitivity and stably produces high-quality images even after 100,000 images are produced.

$$E_{ox1} - E_{ox2} \geq -0.2 \quad (I)$$

On the other hand, when out of the above-mentioned relationship, the bright portion of the resultant photoreceptor is extremely high from the beginning. Therefore, the image

density deteriorates and no image can be produced.

The deterioration of image resolution of the images produced by the comparative photoreceptor 1 in Comparative Example 28 due to repeated use is worse than that of the images produced by the photoreceptor of the present invention because of not including the compound having an alkylamino group effective for oxidizing gases causing blurred images.

Further, the comparative photoreceptor 2 in Comparative Example 29 including only a typical antioxidant in its CTL has a high bright portion potential from the beginning and does not produce good images.

This document claims priority and contains subject matter related to Japanese Patent Application No. 2003-049975 filed on February 26, 2003 incorporated herein by reference.

Having now fully described the invention, it will be apparent to one of ordinary skill in the art that many changes and modifications can be made thereto without departing from the spirit and scope of the invention as set forth therein.